

GenCore version 5.1.3  
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## SUMMARIES

OM protein - nucleic search, using frame\_plus\_p2n model

Run on: October 19, 2002, 07:51:16 : Search time 199 Seconds

(without alignments)  
3951.492 Million cell updates/sec

Title: US-09-807-459-2

Perfect score: 2359

Sequence: 1 MAPSDSVGDTVTKTLAASES.....DPKALIRKVSSTEADNILEK 458

Scoring table:

BLOSUM62  
Xgapop 10.0 , Xgapext 0.5  
Ygapop 10.0 , Ygapext 0.5  
Fgapext 6.0 , Fgapext 7.0  
Delop 6.0 , Delext 7.0

Searched: 1736436 seqs, 858457221 residues

Total number of hits satisfying chosen parameters: 3472872

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Command line parameters:

-MODL=frame\_p2n.model -DEV=xlh  
-O=/cgn2\_1/USPRO.spool/US09807459/runat\_18102002.141111-28571/app-query.fasta.1.647  
-DB=N\_Geneseq\_032802 -QFMT=fastap -SUFFIX=ring -MINMATCH=0.1 -LOOPEXT=0  
-LOOPEXT=0 -UNITS=bits -START=1 -END=-1 -MATRIX=blomsum2 -TRANS=human40.cdi  
-LIST=45 -DOCALLIGN=200 -THR SCORE=pct -THR MAX=100 -THR MIN=0 -ALIGN=40  
-MODE=LOCAL -OUTFMT=ptc -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000  
-USER=US09807459 -GCGN\_1.1\_48 -truncat\_18102002.141111.28571 -NCPU=6 -ICPU=3  
-NO\_XLPRY -NO\_MMAR -LARGEQUERY -NEG\_SCORES=0 -WAIT -LONGLOC -DEV=TIMEOUT=120  
-WARN\_TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -Fgapop=6 -Fgapext=7  
-YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database : N\_Geneseq\_032802:\*

1: /SIDS1/gcgdata/hold-geneseq/geneseqn-emb1/NA1980.DAT:\*  
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22: /SIDS1/gcgdata/hold-geneseq/geneseqn-emb1/NA2001A.DAT:\*  
23: /SIDS1/gcgdata/hold-geneseq/geneseqn-emb1/NA2001B.DAT:\*  
24: /SIDS1/gcgdata/hold-geneseq/geneseqn-emb1/NA2002.DAT:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

Result No.	Score	Query Match	Length	DB ID	Description
1	2359	100.0	1828	22 AAF59961	Babesia caballi me
2	996.5	42.2	1542	14 AAO47074	21B4/rhoptry genes
3	827.5	35.1	1962	16 AAO90252	Babesia merzoiite
4	826.5	35.0	1990	14 AAO33064	Encodes Babesia bo
5	826.5	35.0	1990	17 AAT18995	Babesia merzoiite
6	770.5	32.7	1371	14 AAO47076	B. canis 21B4/rhop
7	759.5	32.2	1491	14 AAO47075	21B4/rhoptry antig
8	161	6.8	131	AAO26065	21B4 gene clone pr
9	147	6.2	1664976	19 AAT21209	Methanococcus jam
10	138	5.8	170	AAO26066	21B4 gene clone pr
11	134	5.7	26776	20 AAX20254	Borrelia burgdorfe
12	122.5	5.2	111309	20 AAX20250	Borrelia burgdorfe
13	122.5	5.2	910715	20 AAX20248	Borrelia burgdorfe
14	121	5.1	11165	21 AAI15186	DNA encoding Esche
15	120	5.1	3883	20 AAV71915	S. cerevisiae C1N8
16	120	5.1	3884	22 AAH78010	Nucleotide sequenc
17	118.5	5.0	4766	15 AAO70102	Malariai P1EMP3 ep
18	118.5	5.0	7326	21 AAO70095	Plasmodium falcipa
19	117	5.0	33303	21 AAO81514	N. meningitidis pa
20	117	5.0	349980	21 AAF21610	Neisseria meningit
21	115	4.9	6637	23 AAS71042	DNA encoding novel
22	113.5	4.8	4443	23 ABL04821	Drosophila melanog
23	113.5	4.8	10478	20 AAV84691	Arabidopsis ESD4 (
24	113	4.8	4282	23 AAS76165	DNA encoding novel
25	113	4.8	11087	23 AAS74637	DNA encoding novel
26	112.5	4.8	2375	22 AAK68993	Human immune/haema
27	112.5	4.8	42738	22 AAK68992	Human immune/haema
28	111	4.7	1422	20 AAX35720	cDNA encoding a pr
29	111	4.7	1613	20 AAX35721	cDNA encoding a pr
30	111	4.7	3278	18 AAT73285	K. lactis origin o
31	111	4.7	3278	18 AAT73285	Kluyveromyces lact
32	110	4.7	1777	11 AAO06842	RNA segment 4 codi
33	109.5	4.6	1816	23 AAS74418	DNA encoding novel
34	109.5	4.6	2277	23 AAS69641	DNA encoding novel
35	109.5	4.6	2277	23 AAS71279	DNA encoding novel
36	109.5	4.6	2277	23 AAS74306	DNA encoding novel
37	109.5	4.6	2277	23 AAS74426	DNA encoding novel
38	109.5	4.6	2277	23 AAS74599	DNA encoding novel
39	109.5	4.6	2277	23 AAS79126	DNA encoding novel
40	109.5	4.6	2394	23 AAS84093	DNA encoding novel
41	109.5	4.6	3569	23 AAS78798	DNA encoding novel
42	109.5	4.6	1664976	19 AAV21209	Methanococcus jam
43	109	4.6	1572	20 AAX61708	B. burgdorferi ant
44	109	4.6	2746	23 AAS82185	DNA encoding novel
45	109	4.6	3165	23 AAS78781	DNA encoding novel

## ALIGNMENTS

RESULT 1  
AAFS9961  
ID AAF59961 standard: CDNA, 1828 BP.

AAFS9961;  
22-MAY-2001 (first entry)

Babesia caballi merzoiite 48 kd rhoptry protein-encoding CDNA.

Merzoiite protein: 48 kd rhoptry protein; antigen; antibody;

recombinant production; diagnosis; equine babesiosis;

parasitic infection; veterinary; ss.

Babesia caballi.

WO200112813-A1.

PD 22-FEB-2001.  
XX  
PF 13-AUG-1999; 99MO-JP04386.  
XX  
PR 13-AUG-1999; 99MO-JP04386.  
XX  
PA (KAGA) CHERO-THERAPEUTIC RES INST.  
PI (MIKA) MIKAMI T.  
XX  
PI Mikami T, Ikeda H, Igarashi I, Suzuki N, Nagasawa H, Fujisaki K;  
DR WPI: 2001-202867/20.  
DR P-PSDB; AAB50669.  
XX  
PT Gene encoding merozoite protein of Babesia caballi for diagnosis of  
XX equine babesiosis caused by this organism -  
PS Claim 3; Page 19-22; 27pp; Japanese.  
XX  
CC The invention relates to a 48 kD merozoite rhoptry protein from Babesia  
CC caballi (AAB50669) and cDNA encoding it (AAF59961). The invention also  
CC relates to phase vectors containing a nucleic acid encoding the  
CC merozoite protein, a method for the recombinant production of the  
CC protein, an antibody against the protein, and a method for the diagnosis  
CC of equine babesiosis from horse blood samples by using the antibody to  
CC detect Babesia caballi merozoites, or by using the 48 kD protein as an  
CC antigen to detect anti-Babesia caballi antibodies. The 48 kD merozoite  
CC protein, or an antibody specific for the protein may be used for the  
CC diagnosis of equine babesiosis caused by Babesia caballi. The present  
CC sequence represents cDNA encoding the Babesia caballi merozoite 48 kD  
CC rhoptry protein.  
XX  
SQ Sequence 1828 BP; 523 A; 412 C; 460 G; 433 T; 0 other;  
Alignment Scores:  
Pred. No.: 1.46e-22 Length: 1828  
Score: 2359.00 Matches: 458  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 100.00% Indels: 0  
DB: Gaps: 0  
US-09-807-459-2 (1-458) x AAF59961 (1-1828)  
QY 1 MetAlaProSerAspSerValGlyAspValThrIleuLeuAlaAlaSerGluSer 20  
DB 39 ATGGCTCCAGCGACTGTGGGAGCTGACTAGACCTTATGGCTCCAGCGAAAGT 98  
QY 21 ValAspSerAlaAlaAsnAlaTyrMetIleAsnSerAspMetSerAspTyrLeuSerAla 40  
DB 99 GTGAGACTCAGCTGCCAATGCTTATATGATCAACAGTACATGACATGACCTTACTTGGCT 158  
QY 41 ValSerAspAsnPheAlaGluArgIleCysSerGlnValProLysGlySerAsnCysSer 60  
DB 159 GTGCTCACAACCTTGGCGAGCGCATTTGACGTACAGGCTCCCTAAGGAGAGTAACTGCACT 218  
QY 61 AlaSerValSerAlaTyrMetSerArgCysAlaLysGlnAspCysLeuThrLeuGlnSer 80  
DB 219 GCTTCGCTTACGCCATGACATGAGCTGCTGGCTCAACAGAGCTCCGATGCTCTCAAAAGT 278  
QY 81 LeuLysTyrProLeuGlnAlaLysTyrGlnProLeuThrLeuProAspProTyrGlnLeu 100  
DB 279 CTTAAAGTACCTCTTGAAGCTAAAGTACCAACCCCTGACCTTCTGACCCCTTACAGTTTG 338  
QY 101 GluAlaAlaPheIleLeuPheLysGluSerAspAlaAsnProAlaAsnSerThrGluLys 120  
DB 339 GAGGCCCATTTATACCTTTCAGAGAGAGTACCTAATCCGCGCAATAGACTGAAGAAG 398  
QY 121 ArgPheThrPheAlaArgPheArgGlyLysAsnHisSerTyrPheHisAspLeuValPhe 140  
DB 399 CGGTTTCGATGCGCTTTCAGAGAGGCAAGACACAGTACTTCCACGACTTGTCTTC 458  
QY 141 AsnLeuLeuGluLysAsnValThrArgAspAlaAspAlaThrAspIleGluAsnPheAla 160

DB 459 AATCTGCTGGAGAAACGTACTCGACCGCATGCTACTGACATTCGAACTTGGCG 518  
QY 161 SerArgTyrLeuTyrMetAlaThrLeuTyrTyrIleuThrAsnValAspGluPhe 180  
DB 519 TCCAGTACCTCTGACATGCGCACCTTTACTACAAAGCATACAGCAATGTGATGACTTC 578  
QY 181 GlyAlaSerPhePheAsnLysLeuSerPheThrGlyLeuPheGlyTyrPglyLys 200  
DB 579 GGTGCTAGCTCTTAAACAAGTGTCTTTCACACTGCTGGGTGTTGGCTGGGCGATCAAG 638  
QY 201 ArgAlaLeuLysGlnIleIleArgSerAsnLeuProLeuAspIleGlyThrGlnHisSer 220  
DB 639 AGGCACCTTAAAGCATTTATTCGCTTAACTGCCCTTGACATCGGACAGAACAGC 698  
QY 221 ValSerArgLeuGlnHisIleThrSerSerTyrLysAspTyrMetAspThrGlnIlePro 240  
DB 699 GTCAAGTGGCTGACAGCATTTACGACGATTACAGAGATTACATGATACGACATTCCT 758  
QY 241 AlaLeuProLysPheAlaLysArgPheSerLeuMetValGlnArgLeuAlaThr 260  
DB 759 GCATCGCCCAAGTTTGGCAACGTTTCTCCTTATGATGACGAGAGCTGCGCACCC 818  
QY 261 ValIleGlyTyrValAspThrProTyrTyrLysTyrTyrMetLysLeuLysAsnPhe 280  
DB 819 GTGCTGCTTACGTGACACCCCTGGTATMAAGTGTACATGAAAGCTGAAACACTTT 878  
QY 281 MetValAsnArgValPheIleProThrLysPhePheAsnLysGluIleArgGluPro 300  
DB 879 ATGGTGAACGGGTGTTTCATTCACAAAGATTTCTCATTAAGAAATTCGAGACCT 938  
QY 301 SerLysAlaLeuLysGluLysValSerThrAspThrLysAspLeuPheGluAsnLysIle 320  
DB 939 AGTAAGGATTTAAAGAAAGGTGTCAACCGACACCAAGATTTATTCGAGAACAAATTT 998  
QY 321 GlyGlnGlyThrValAspPhePheAsnLysGluIleArgAspProSerLysAlaLeuLys 340  
DB 999 GGGCAGGCTACTGTGCACTTCTCAATAGGAAATTCGTGACCTTATAGGCAATTAAAA 1058  
QY 341 GluLysValSerAsnAspAlaLysAspLeuPheGluAsnLysIleGlyGlnGlyThrVal 360  
DB 1059 GAANAAGTGTCAACGACGCCCAAGATTTATTCAGAACACAAATTTGGCAGGCTACTGTG 1118  
QY 361 AspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLysValSerThr 380  
DB 1119 GACTTCATCATTAACGAATTCGTGACCTGTAGGCAATTAAAGAAAGTGTCAACG 1178  
QY 381 GlyValGluAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPheIleAsnAsn 400  
DB 1179 GGGCCGAGGATTTTATTCGAAACAAATTTGGGAGGCTACTGTGACTTCATCAATAAC 1238  
QY 401 GluIleArgAspProSerLysAlaLeuIleArgLysValTyrThrGluAlaAspLeu 420  
DB 1239 GAATTTGCTGACCTTACTAGGCACTTAAATTAAGAAAGTGTACACCGGCGCATGATTTA 1298  
QY 421 PheGluAsnLysIleGlyGlnGlyThrValAspPheIleAsnLysGluIleArgAspPro 440  
DB 1299 TTGAGAAACAAATTTGGGAGGCTAGTGTGACCTTCAATCAATAGAAATTCGTGACCT 1358  
QY 441 SerLysAlaLeuIleArgLysValSerThrGluAlaAsnLeuLeuGluLys 458  
DB 1359 AGTAAGCATTTAAATMAAGAAAGTGTACCGAGGCGCATTAATTTATTTGAGAAA 1412  
RESULT 2  
ID AA047074 standard; DNA: 1542 BP.  
XX  
AC AA047074;  
XX  
DT 13-JAN-1994 (first entry)  
XX  
DE 21B4/rhoptry genes 1-4 representative DNA.  
XX



Db 1369 GGTGCCCATTTAAAGA-ATCATTTGAGAACTTGGACACAAAGCG 1412  
RESULT 3  
AAQ90252  
ID AAQ90252 standard; cDNA: 1962 BP.  
XX  
AC AAQ90252;  
XX  
DT 14-NOV-1995 (first entry)  
XX  
DE Babesia merozoitae p58 cDNA.  
XX  
KM Merozoite; surface protein; antigen; p58; babesiosis; vaccine; ss.  
OS Babesia bigemina.  
XX  
FH Key Location/Qualifiers  
FT CDS 186..1628  
FT /\*tag- a  
FT sig\_peptide 186..248  
FT /\*tag- b  
FT mat\_peptide 249..1631  
FT /\*tag- c  
FT polyA\_signal 1884..1889  
FT /\*tag- d  
XX  
PN US542428-A.  
XX  
PD 06-JUN-1995.  
XX  
PF 27-MAR-1987; 87US-0031328.  
XX  
PR 06-DEC-1991; 91US-0803636.  
PR 27-MAR-1987; 87US-0031328.  
PR 01-MAR-1991; 91US-0663255.  
XX  
PA (UNIW ) UNIV WASHINGTON STATE.  
XX  
PI Davis WC, McElwain TF, McGuire TC, Perryman LE;  
XX  
DR MPI: 1995-214706/28.  
DR P-PSDB; AAR77249.  
XX  
PT Babesia merozoitae 45 kD surface protein from B. bigemina - used in  
XX vaccines for the prophylaxis of bovine babesiosis.  
XX  
PS Disclosure: Column 31-34; 30pp: English.  
XX  
CC Antigenic surface proteins (45, 55 and 58 kDa) were isolated from  
CC the intracythrotic merozoite stage of B. bigemina JG-29. The 58  
CC kDa surface protein (AAR77249) was characterized, and encoding  
CC cDNA (AAQ90252) was isolated from a lambda GEM1 library.  
XX  
SO Sequence 1962 BP: 506 A; 442 C; 492 G; 522 T; 0 other:  
SO  
Alignment Scores:  
Pred. No.: 1e-71 Length: 1962  
Score: 827.50 Matches: 181  
Percent Similarity: 59.12% Conservative: 75  
Best Local Similarity: 41.80% Mismatches: 140  
Query Match: 35.08% Indels: 37  
DB: 16 Gaps: 12  
US-09-807-459-2 (1-458) x AAQ90252 (1-1962)  
Qy 4 SerAspSerValGlyAspValThrLysThrLeuAlaAlaSerGluSerValAspSer 23  
Db 282 GAGAGGGGTGGAGATGTCTCAAGACCTTGTGGAAGCCAAATGAGGTTTCAATGCT 341  
Qy 24 AlaAlaAsnAlaTyrMetIleAsnSerAspMetSerAspTyrLeuSerAlaValSerAsp 43  
Db 342 GAATGGAAGCAACATCAGCTCAACAAGATATGCAAAAGCATTTGTCTAATGTTAAGAG 401

Qy 44 AsnPhaAlaGluArgIleCysSerGlnValProLysGlySerAsnCysSerAlaSerVal 63  
Db 402 ACCATTGTTGGAGAGTCTCCGCAAGAAAGTGTCTGGAACCTTACCTGGCGTGAAGCGTA 461  
Qy 64 SerAlaTyrMetSerArgCysAlaLysGlnAspCysLeuThrLeuGlnSerLeuLysTyr 83  
Db 462 ATTGCCATATGTAACCGTTGTGATGAGCGCATTTGTCTGACGCTTGACAGCATG----- 515  
Qy 84 ProLeuGluAlaLysTyrGlnProLeuThrLeuProAspProTyrGlnLeuGlnAla 103  
Db 516 -----AAGTACAAAGCGTTGAGTCTGCAAAATCTTACCAAGTTGAGCGCTGCC 563  
Qy 104 PheIleLeuPheLysGluSerAspAlaAsnProAlaAsnSerThrGluLysArgPheTyr 123  
Db 564 TTCATGCTTTTACGGAAGATGATCTTAACCTCGCAAGAATGAGTGAAGCGCTTCTGG 623  
Qy 124 MetArgPheArgArgGlyLysAsnHisSerTyrPheHisAspLeuValPheAsnLeu 143  
Db 624 ATGCGTTGAGG-----AGCAGCCAGCGGCACTACCATCTTGTGTGCTTGTG 677  
Qy 144 GluLysAsnValThrArgAspAlaAspAlaThrAspIleGluAsnPhaAlaSerArgTyr 163  
Db 678 AAGAAGAAATGTTGACGCGACCTTAATCCAATGATGTTGAGAACCTTGCCATCGCATAC 737  
Qy 164 LeuTyrMetAlaThrLeuTyrTyrLysThrTyrThrAsnValAspGluPheGlyAlaSer 183  
Db 738 TTCTACATGACATGATGTTGATACAAAGACTTACGCGGTTGACTTAAAGCGGCTAAG 797  
Qy 184 PhePheAsnLysLeuSerPheThrThrGlyLeuPheGlyTyrGlyIleLysArgAlaLeu 203  
Db 798 TTCTTCAACACACCTTCTTCAAACTCGCTTGTGTTGTTGATTCAGAAAGCGTTG 857  
Qy 204 LysGlnIleIleArgSerAsnLeuProLeuAspIleGlyThrGluHisSerValSerArg 223  
Db 858 AAGCGTTGGTTAGAGACACTTCCCGTTGACCTTGGAAC-----CACCGTAGGCCAC 914  
Qy 224 LeuGlnHisIleThrSerSerTyrLysAspTyrMetAspThrGlnIleProAlaLeuPro 243  
Db 915 ATCCGCGAAATGATGATGAGCGGCTACGCGAGTACATGATGACCCAGCTCCGATGACC 974  
Qy 244 LysPheAlaLysArgPheSerLeuMetValValGlnArgLeuAlaThrValAlaGly 263  
Db 975 TCGTTCGCTGACGCTTCTTCCAAAGATGGCTACTAGACTCTGTGTTACCTCCGAC 1034  
Qy 264 TyrValAspThrProTyrTyrLysTyrTyrMetLysLeuLysAsnPhaMetValAsn 283  
Db 1035 TACGTCATTTGCCCGCGGTACAAAGAGGTGTCACAGAACTTCAAGGAATTCATTGTGAC 1094  
Qy 284 ArgValPheIleProThrLysLysPheAsnLysGluIleArgLupProSerLys--- 302  
Db 1095 ---TCTTATACGACCTCGCCCAAGTTGATTAAGACGACGTCTCAGCCTTAAGACT 1151  
Qy 303 AlaLeuLysGluLysValSerThrAspThrLysAspLeuPheGluAsnLysIleGln 322  
Db 1152 GCTTACACAAAGCTGCTCCCGACAGACAGCGAGCTATCAGGAATGCTCGCTGTA 1211  
Qy 323 GlyThrValAspPhePheAsnLysGluIleArgAspProSerLysAlaLeuLysGluLys 342  
Db 1212 AGCACCAAGCATATTTGCCAAC---GGTGTAGCTGATTTGTCAAGATGATTAAAGAG--- 1265  
Qy 343 ValSerAsnAspAlaLysAspLeuPheGluAsnLysIle-----Gly 356  
Db 1266 -----CCTAAGCCAAACAAATTAATTCGTGAGAGCTGCTCACTTACTTCTAAGCA 1316  
Qy 357 GlnGlyThrValAspPheIleAsnAsnGluIleArgAsp-----ProSerLysAlaLeu 374  
Db 1317 AAGGAGCGGTTGACGACGCTGTTAAGAAGGTTAATCCGTTGCTCCGATTAAG----- 1370  
Qy 375 IleArgLysValSerThrGlyAlaGluLysPhePheGluAsnLysIleGlyGlnGlyThr 394  
Db 1371 ---CAAAAGGGCGGACCAACATCCGAGACAGCTGTAGAGAAACGTTCCGCTGCG--- 1424

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Oy 395 ValAspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLysValTyr 414
    ||| ||| ||| |||
Db 1425 ---GATTCGCGGAACGTAATTTGAGTCCCTCGAAGAACAAATCGATCGTGTACT 1481
Oy 415 Thr-----GluAlaAspAsp 419
    ||| ||| ||| |||
Db 1482 ACTCAGAGAGTTAACAGCAGAGAGAGTGTATGCCGACGAT 1520

RESULT 4
AAO33064
ID AAO33064 standard; DNA; 1990 BP.
XX
XX AAO33064;
XX
XX 06-MAY-1993 (first entry)
XX
XX Encodes Babesia bovis 60kD immunoreactive merozoite surface epitope.
XX
XX babesiosis; cows; cattle; bos taurus; babesia bovis; babesia bigemina;
XX merozoite; schizont; ss.
XX
XX Babesia bovis.
XX
XX Key Location/Qualifiers
XX CDS 1..1990
XX FT /*tag= a
XX
XX PN US5171685-A.
XX
XX PD 15-DEC-1992.
XX
XX PE 04-APR-1990; 90US-0504461.
XX
XX PR 04-APR-1990; 90US-0504461.
XX
XX (USFL ) UNIV FLORIDA.
XX (USDA ) US SEC OF AGRIC.
XX
XX Davis WC, Goff WL, Hines SA, Jasmer DP, McElwain TF;
XX Mcgillire TC, Palmergh, Perryman LE, Reduker DW;
XX
XX WPI: 1993-008582/01.
XX P-PSDB; AAR30613.
XX
XX DNA encoding Babesia bovis protein - is used as probes and for
XX prodn. of polypeptide(s) for use in vaccines and for prodn. of
XX antibodies
XX
XX Example 19; Fig 3; 20pp; English.
XX
XX This sequence encodes an immunoreactive epitope located on the
XX surface of babesia bovis merozoites. A. B. bovis cDNA expression
XX library was constructed using poly A(+) RNA isolated from B. bovis
XX infected blood cultures. Plaques were immunoscreened using rabbit
XX anti-Bv60 sera, and positive plaques tested for reactivity with
XX monoclonal antibodies that recognised a Bv42 surface exposed epitope
XX as well as an isotype control monoclonal antibody and normal rabbit
XX serum. Lambda rBv60 phagemid DNA was isolated from bacteria, and
XX then restriction digested.
XX
XX Sequence 1990 BP; 628 A; 437 C; 398 G; 527 T; 0 other;
XX
Alignment Scores:
Pred. No.: 1,28e-71 Length: 1990
Score: 826.50 Matches: 178
Percent Similarity: 51.30% Conservative: 79
Best Local Similarity: 35.53% Mismatches: 195
Query Match: 35.04% Indels: 49
Db: 14 Gaps: 6
US-09-807-459-2 (1-458) x AAO33064 (1-1990)
Oy 1 MetaIaProSerAspSerValGlyAspValThrLysThrLeuLeuAlaIaSerGluSer 20

```

```

Db 212 CTCGCTCCAGCTGAAGTGTAGTGTATTTAACTCCACATTTGGAACAGCTGATCTTG 271
Oy 21 ValAspSerAlaIaIaAsnAlaTyrMetIleAsnSerAspMetSerAspTyrLeuSerAla 40
    ||| ||| ||| |||
Db 272 ATGACTCTCCGTGACGACATGACCAACATTACTAGAGATATGAAACATGTTTGGACAAAT 331
Oy 41 ValSerAspAsnPhaIleArgIleCysSerGlnValProLysGlySerAsnCysSer 60
    ||| ||| ||| |||
Db 332 GGTGCGAGCAGATGTGTAATGATGTTTGTCTTAATGCTCTGAGACCTCAACTGCTGCT 391
Oy 61 AlaSerValSerAlaTyrMetSerArgCysAlaLysGlnAspCysLeuThrLeuGlnSer 80
    ||| ||| ||| |||
Db 392 GAGGTAGTTTAACTATATGCTGACCGTTGTAATGATGACGATGCTTACGATTTGACAAAT 451
Oy 81 LeuLysTyrProLeuGluAlaLysTyrGlnProLeuThrLeuProAspProTyrGlnLeu 100
    ||| ||| ||| |||
Db 452 GTCAAAATATCCGTTGTATCAAGAGTACCAACCTCTATCTCTCCAAACCTTACCGATTG 511
Oy 101 GluAlaIaPheIleLeuPheLysGluSerAspAlaAsnProIaAsnSerThrGlnLys 120
    ||| ||| ||| |||
Db 512 GATGCTGCGCTTCAGATTTGTTCAAGAGAGTGCATCCGACCTGCCAAGAACACCGTAAAT 571
Oy 121 ArgPheTyrMetArgPheArgArgGlyLysAsnHisSerTyrPheHisAspLeuValPhe 140
    ||| ||| ||| |||
Db 572 CGCGAATGGTTGGCTTACAAATGAGCGAACCATGGTATACCACTACTTCGTCACAT 631
Oy 141 AsnLeuLeuGluLysAsnValThrArgAspAlaIaThrAspIleGluAsnPhaIa 160
    ||| ||| ||| |||
Db 632 GGTCTGTGTGAACAACATGTTGTGCACGAGGAAGAACTCCGATGTGAATATCTTGTTC 691
Oy 161 SerArgTyrLeuTyrMetAlaThrLeuTyrTyrLysThrTyrThrAsnValAspGluPhe 180
    ||| ||| ||| |||
Db 692 AACAGGACTCTATATGCTGCTACCATGAACTACAGACACTATTTGACAGTAACAGATATG 751
Oy 181 GlyAlaSerPhePheAsnLysLeuSerPheThrThrGlyLeuPheGlyTyrGlyLys 200
    ||| ||| ||| |||
Db 752 AACGCCAAGTTTCAACAGATTCAGCTTCATCAACAAAGATTTAGTCTGATATAG 811
Oy 201 ArgAlaLeuLysGlnIleIleArgSerAsnLeuProLeuAspIleGlyThrGlnHisSer 220
    ||| ||| ||| |||
Db 812 CAACATGTGATGATATCAACAGTGAAGTGAATGTTCCGAAAGATTTT---GAAGAAAGAGC 868
Oy 221 ValSerArgLeuGlnHisIleThrSerSerTyrLysAspTyrMetAspThrGlnIlePro 240
    ||| ||| ||| |||
Db 869 ATCGAAGATATCACTCACTACTACTAGCAGCTACGAGAAATATCTTGTGACCCAGATTCCA 928
Oy 241 AlaLeuProLysPheAlaLysArgPheSerLeuMetValValGlnArgLeuLeuAlaThr 260
    ||| ||| ||| |||
Db 929 ACTCTTTCCAAGTTTGACGCTGCTATGCTGACATGCTGAAGAAAGTTCGTCGCGTAGC 988
Oy 261 ValAlaGlyTyrValAspThrProTyrTyrLysTyrTyrMetLysLeuLysAsnPha 280
    ||| ||| ||| |||
Db 989 TTGACTCGACCTTGTAGAGCTCTTGTGTCAAAAGATGATAAAGAAATTCAGAGACTT 1048
Oy 281 MetValAsnArgValPheIleProThrLysLysPhe----- 292
    ||| ||| ||| |||
Db 1049 TTCTTAATAAAGCTTACCCACACTACAAAGAGATTCATCGAGCATCTAAAGAAATTACC 1108
Oy 292 ----- 292
Db 1109 AAAAATCTGTAAGCAATGTTGCTGAGCCCACTAAAGTTTATGACGACACTCAC 1168
Oy 293 -----PheAsnLysGluIleArgGluProSerLysAlaLeuLysGlu 306
    ||| ||| ||| |||
Db 1169 GAAAAAACCAAGCTATCTGAAGAGAAATGTAGCCGAACTCTAAGACTTTTTCACAG 1228
Oy 307 LysValSerThrAspThrLysAspLeuPheGluAsnLysIleGlyGlnGlyThrValAsp 326
    ||| ||| ||| |||
Db 1229 GAGGCTCTCACTCACTCAACCACTTCTTGATGAGAACATTTGCCAACCCCAAGAGAG 1288
Oy 327 PhePheAsnLysGluIleArgAspProSerLysAlaLeuLysGluLysValSerAsnAsp 346
    ||| ||| ||| |||

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Db 1289 TTTTTCAGGAGAGCTCCCAAGCCACTAAACATTTCTAGACGAAACATCGTCACCA 1348  
 QY 347 AAlaysAspLeuPheGluAsnLysIleGlyInglYThrValAspPheIleAsnGlu 366  
 Db 1349 ACCAAGGAGTTCTTC---AGGAGGCTCTCTCAAGCCACTAAGCACTCTTGGCGAGAT 1405  
 QY 367 IleArgAspProSerLysAlaLeuIleArgLysValSerThrGlyAlaGluAspLeuPhe 386  
 Db 1406 ATTGCTCAACTACTAAAGAAATTTTTCAGAGATGTCCCTCAAGTACCAAGAGTTTATA 1465  
 QY 387 GluAsnLysIleGlyInglYThrValAspPhe----- 397  
 Db 1466 ACTGAGAACATTGCTCAACCACTAGAGATGTCGAGAGAGAGTTCTTCATGCTTACCATG 1525  
 QY 398 -----IleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLysValThr 415  
 Db 1526 AAGTCTTGATGAAGAACATTGCTCAACCTGCGAAGAAATCATATGAGTTTGCTACA 1585  
 QY 416 GluAlaAspAspLeuPheGluAsnLysIleGlyInglYThrValAspPheIleAsnLys 435  
 Db 1586 GCGCGCAAGAAAT---TTCATTTCCGACGCCATGAGAGTACTAAGCACTTTTAAACGA 1642  
 QY 436 GluIleArgAspProSerLysAlaLeuIleArg---LysValSerThrGluAlaAspAsn 454  
 Db 1643 ACTGTGGCCCACTTACCAAGGAATTCCTGAACGAGCTTTAGAAACTACTAAAGACGCA 1702  
 QY 455 Leu 455  
 Db 1703 TTA 1705  
 RESULT 5  
 AAT18995  
 ID AAT18995 standard; cDNA; 1990 BP.  
 XX AAT18995:  
 AC AAT18995:  
 XX  
 DT 15-OCT-1996 (first entry)  
 DE Babesia merozoitae surface protein cDNA clone Bv60.  
 XX  
 KM Babesia merozoitae protein; vaccine; probe; diagnosis; ss.  
 XX  
 OS Babesia bovis.  
 XX  
 FH Key Location/Qualifiers  
 FT CDS 122..1819  
 FT /\*tag= a  
 XX  
 FN US518916-A.  
 XX  
 PD 21-MAY-1996.  
 XX  
 PF 04-APR-1989; 89US-0333155.  
 XX  
 PR 04-APR-1990; 90US-0504461.  
 PR 04-APR-1989; 89US-0333155.  
 PR 14-DEC-1992; 92US-0989616.  
 PR 21-NOV-1994; 94US-0342480.  
 XX  
 PA (USDA ) US SEC OF AGRIC.  
 XX  
 PI Goff WL, Jasmer DP, McElwain TF, McGuire TC, Reduker DW;  
 PI Stillier D;  
 XX  
 DR WPI: 1996-259067/26.  
 DR  
 DR P-PSDB: AAR97981.  
 XX  
 XX New fragment of Babesia bovis genomic DNA - useful as a probe for  
 PT detecting Babesia infection  
 XX  
 PS Example 18; Fig 3; 19pp; English.  
 CC A cDNA clone (AAT18995) codes for Bv60, a 60 kDa immunoreactive

CC protein (AAR97981) located on the surface of Babesia bovis merozoites.  
 CC It was isolated from a blood-stage B. bovis cDNA library in lambda  
 CC ZAPRI by subcloning into Bluescript SK(-) and immunoscreening using  
 CC monospecific anti-Bv60 antiserum. Bv60, Bv4 and Bv42 (see also  
 CC AAT18993 and AAT18994) DNA sequences can be used to make recombinant  
 CC proteins useful as vaccines for the prophylaxis of bovine babesiosis.  
 CC They can also be used as diagnostic probes.  
 CC  
 XX

SQ Sequence 1990 BP; 628 A; 437 C; 398 G; 527 T; 0 other.

#### Alignment Scores:

Pred. No.:	1,28e-71	Length:	1990
Score:	826.50	Matches:	178
Percent Similarity:	51.30%	Conservative:	79
Best Local Similarity:	35.53%	Mismatches:	195
Query Match:	35.04%	Indels:	49
DB:	17	Gaps:	6

US-09-807-459-2 (1-458) x AAT18995 (1-1990)

QY 1 MetaIAspProSerAspSerValGlyAspValThrLysThrLeuLeuAlaSerGluSer 20  
 Db 212 CTCGCTCCAGCTGAGGTGAGTGTAGCTATTTAACTTCACTTGGAAACAGCTGATCTTG 271  
 QY 21 ValAspSerAlaAlaAsnAlaLysMetIleAsnSerAspMetSerAspPyrLeuSerAla 40  
 Db 272 ATGACTCTCCGTGACCACTGACCAACATTTACTAGATATGAAACATGTTTGGCAAT 331  
 QY 41 ValSerAspAsnPhaAlaGluArgLecysSerGlnVal----- 60  
 Db 332 GGTCTGAGACAGATTGTAATGATGTTGCTTAATGCTCTGAGAGACTCCACAGCTGCT 391  
 QY 61 AlaSerValSerAlaLysMetSerArgCysAlaLysGlnAspCysLeuThrLeuGlnSer 80  
 Db 392 GAGTAGATTAACTTATGCTGACCGCTTGTAATGTAAGATGCTTACGATTGACAT 451  
 QY 81 LeuLysTYRProLeuGluAlaLysTYRcInProLeuThrLeuProAspProTyrGlnLeu 100  
 Db 452 GTCAAAATATCCGTTGTATCAAGAGTACCAACCTCTATCTCTCCAAACCTTACCACTTG 511  
 QY 101 GluAlaAlaPheIleLeuPheLysGluSerAspAlaAsnProAlaAsnSerThrGluLys 120  
 Db 512 GATGCTGCTTACATTTGTTCAAGAGAGTGCATGGAACCTCGCCACAGACGCTTAAA 571  
 QY 121 ArgPheTrpMetArgPheArgArgGlyLysAsnHisSerTYRPhaHisAspLeuValPhe 140  
 Db 572 CGCGAATGCTGCTTACAGAAATGAGACGAACATGATTTACACTACTGCTGACT 631  
 QY 141 AsnLeuLeuGluLysAsnValThrArgAspAlaAspAlaThrAspIleGluAsnPhaAla 160  
 Db 632 GGTCTGTTGAAACACAAATGTTGTGCACGACGAAAGCACTACCGATGTAATCTTGTCT 691  
 QY 161 SerArgTYRLeuTYRMetAlaThrLeuTYRTrpTYRTrpAsnValAspLuphe 180  
 Db 692 AACCAAGTACTATATATGCTTACATGAACTACAGACTATTTGACAGTAACGTAATG 751  
 QY 181 GlyAlaSerPhePheAsnLysLeuSerPheThrThrGlyLeuPheGlyTYRdLysIleLys 200  
 Db 752 AACGCCAGTCTTCAACAGATTCAGCTTCACTCACTAAGAGATATTGACGTCGATTTAG 811  
 QY 201 ArgAlaLeuLysGlnIleIleArgSerAsnLeuProLeuAspIleGlyTYRGlHisSer 220  
 Db 812 CAACATTTAGATGATATGATCAGGTGAGGATGCTCTCAAGATTTT---CAAGAAAGAGC 868  
 QY 221 ValSerArgLeuGlnHisIleThrSerSerTYRTrpLysAspTYRMetAspThrGlnIlePro 240  
 Db 869 ATCGAACCTATTCATCACTTACTAGACGTACGAGAAATTCATGCTGACCCAGATTTCA 928  
 QY 241 AlaLeuProLysPheAlaLysArgPheSerLeuMetValValGlnArgLeuLeuAlaThr 260  
 Db 929 ACTGTTCCAAAGTTTGACGCTGTTATGCTGACATGGTGAAGAAAGTCTGCTGCTGATGC 988  
 QY 261 ValAlaGlyTYRValAspThrProTYRTrpTYRTrpLysTYRTrpMetLysLeuLysAsnPha 280

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Db 989 TTGACCTGCTAGCTGAGCTCTTGGTACAAAGATGATTAAGAAATTGAGACTTT 1048
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 281 MetValAsnArgValPheIleProThrLysLysPhe----- 292
      ||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 1049 TTCCTAAAGCGTTACCAACTACCAAGAGTTTCATGAGAGTACTAAGCAAGTTACC 1108
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 292 ----- 292
Db 1109 AAAAATCTCTGAAGCCATCTTCTGAGCCCACTAAAGTTTATGACAGACACTGC 1168
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 293 -----PheAsnLysGluIleArgGluProSerLysAlaLeuLysGlu 306
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 1169 GAAAAAACCAAGGCTATCTGAAAGATGATGAGCAAGCAACTACTAAGACTTTTTCAG 1228
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 307 LysValSerThrAspThrLysAspLeuPheGluAsnLysIleGlyGlnGlyThrValAsp 326
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 1229 GAGGCTCCTCAAGTCAACCAACTCTTCGATGAGAACATTGGCCCAACCCCAAGAG 1288
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 327 PhePheAsnLysGluIleArgAspProSerLysAlaLeuLysGluLysValSerAsnAsp 346
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 1289 TTTTTCAGGGAAGCTCCCAACCCACTAAACATTCTTCTAGACGAAACATCGTCAACCA 1348
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 347 AlaLysAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPheIleAsnAsnGlu 366
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 1349 ACCAAGGAGTCTTC---AGGAGAGCTCTCTCAAGCCACTAAGCACTCTCTAGCGCAAGAT 1405
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 367 IleArgAspProSerLysAlaLeuIleArgLysValSerThrGlyAlaGluAspLeuPhe 386
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 1406 ATTGCTCAACTACTAAAGATTTTCAAGATGTCCCTCAAGTCAACCAAGAGTTATA 1465
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 387 GluAsnLysIleGlyGlnGlyThrValAspPhe----- 397
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 1466 ACTGGAACATTGCTCAACCACTAAGAGTTCCGAGAGGAGGTTCTCATCCTACCATG 1525
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 398 -----IleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLysValThr 415
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 1526 AAAGTCTGAATGAAGAAATTCCTCACTGCAAGCAAGAAATCATGATGATTTGGTACA 1585
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 416 GluAlaAspAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPheIleAsnLys 435
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 1586 GCGCCCAAGAAAT--TTCATTTCCCGACGCCATGAGAGGTACTAAGCAAGTTCTTAAACGAA 1642
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 436 GluIleArgAspProSerLysAlaLeuIleArg--LysValSerThrGluLysAsn 454
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 1643 ACTGTGGCCCAACTACAAAGAAATTCCTGAACGAGCTTTAGAACTACTAAGAGCGCA 1702
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 455 Leu 455
      |||
Db 1703 TTA 1705

RESULT 6
AA047076
ID AA047076 standard; DNA; 1371 BP.
XX
AC AA047076;
XX
DE 13-JAN-1994 (first entry)
XX
DE B. canis 21B4/rhoptry antigen gene 2 DNA.
XX
KW Polymerase chain reaction; PCR; amplify; primer; detection;
KW babesiosis; parasite; Babesia canis; 21B4/rhoptry; antigen; gene;
KW repeat region; immune response; vaccine; ss.
OS Babesia canis.
OS
PN WO9314204-A.
PN
PD 22-JUL-1993.
PD
XX 15-JAN-1993; 93WO-A000012.
XX

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PR 15-JAN-1992; 92AU-0000399.
XX
PA (CSTR ) COMMONWEALTH SCI & IND RES ORG.
XX
PI Dalrymple BP, Peters JM;
XX
DR WPI; 1993-243219/30.
DR P-PSDB; AAR39902.
XX
PT Detecting closely linked gene copies which encode protective
PT antigen against babesiosis - by screening babesial genomic DNA
PT library with oligo-nucleotide probe based partial sequencing of
PT protective antigen and identifying positive clones
PS
XX Claim 24; Fig 7; 55pp; English.
XX
CC This sequence represents the Babesia canis 21B4/rhoptry antigen gene
CC 2. This sequence was determined from restriction fragments from the
CC clone B. canis lambda GBM-11 #9. B. canis was found to contain two
CC genes which are related to the B. bovis 21B4 gene. Gene 1 and gene 2
CC are very similar but gene 2 appears to contain a large number of
CC repeats. Babesia antigen genes can be used in the production of a
CC combined vaccine which will stimulate a greater immune response and
CC afford broader immunity than a single antigen vaccine. See also
CC AA047068-74.
XX
SQ Sequence 1371 BP; 418 A; 312 C; 325 G; 318 T; 0 other;

Alignment Scores:
Pred. No.: 2,556-66 Length: 1371
Score: 770.50 Matches: 161
Percent Similarity: 55.65% Conservative: 95
Best Local Similarity: 35.00% Mismatches: 161
Query Match: 32.66% Indels: 43
DB: 14 Gaps: 9

US-09-807-459-2 (1-458) x AA047076 (1-1371)
Qy 1 MetAlaProSerAspSerValIleGlyAspValThrLysThrLeuLeuAlaIleSerGluSer 20
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 91 CTTTCTAAATCAGATGAGAGCCGAAAGACCTTGCTCTCACTCTTCACTGACGCATCG 150
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 21 ValAspSerAlaIleAsnAlaThrMetIleAsnSerAspMetSerAspThrLeuSerAla 40
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 151 ACGAGAGCGCTTTACAGAGCTACAGAAATGATGCTGCTATGCAAACTTTCAACCGT 210
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 41 ValSerAspAsnPheAlaGluArgIleCysSerGlnValProLysGlySerAsnCysSer 60
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 211 CCGAGGAAAGAGGAGAGAGAGCTGCTGTGGAACATCGCAGAGAGAGACTGAATGTGAG 270
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 61 AlaSerValSerAlaThrMetSerArgCysAlaIleGlnAspCysLeuThrLeuGlnSer 80
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 271 AAGAGCGTAGCGAGTATGTTGAAGACTCGTCAGGTACGACTGCTTGTGCAATGGAAC 330
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 81 LeuLysTyProLeuGluAlaLysTyArgLysProLeuThrLeuProAspProTyGlnLeu 100
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 331 CAGAACTACCTCAGAGAGAAAGATACAGCCGCTACCCCTTCCCAACCTTATCATATG 390
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 101 GluAlaIlePheIleLeuPheLysGluSerAspAlaAsnProAlaIleAsnSerThrGluLys 120
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 391 GAGCCCGCTTCTATGCTTTCAGAAACAGTGAATCAAACTTAATAAACCAACCAAGGAA 450
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 121 ArgPheThrMetArgPheArgArgGlyLysAsnHisSerTyPheHisAspLeuValPhe 140
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 451 GCTTCTGGATCGTTTCTGTCACGCGGAGCTTATGCGCCCTATTCACATTTCCGGTG 510
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 141 AsnLeuLeuGluLysAsnValThrArgAspAlaAspAlaThrAspIleGluAsnPheAla 160
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 511 AATATTCTATATAAAACCTCAGCGATACATGCGATGATTAACCTCAGAGTTTCGTA 570
      ||||| ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Qy 161 SerArgTyLeuTyMetAlaThrLeuTyTyLysThyTyThrAsnValAspGluPhe 180
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::
Db 571 CGCAATATGCTTACATGCGCACCATGTATATACAGACATACACCGCTTGTGATGTGTA 630
      ::::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||::: |||||:::

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Oy	181	GLVALASerPhePheAsnLysSerPheThrGlyLeuPheGlyProGlyLys	200
Db	631	AATGCCAGAGATCATTAACAAAATTGCTTCTCCCGCACTGTTCGGAAGCAGATTAG	690
Oy	201	ArgAlaLeuLysGlnIleIleArgSerAsnLeuProLeuAspIleGlyThrGlnHisSer	220
Db	691	AACCCCTACAGAAATATAATCCGAGTAACATTCGGAAGACTTGCGC---AAGTACAC	747
Oy	221	ValSerArgLeuGlnHisIleThrSerSerTyrLysAspTyrMetLysPheThrGlnPro	240
Db	748	GTTACCGCCGAGAGCCAGATGGGTGGCTACGAGAGTCACTGTATGAAGCAAGTACCA	807
Oy	241	AlaLeuProLysPheAlaLysArgPheSerLeuMetValGlnArgLeuAlaThr	260
Db	808	TCTCTCCAAATTTCCGCAAGAAATACGCCGGAATGTGTGAAGAGTTTAATTAAGAT	867
Oy	261	ValAlaGlyTyrValAspThrProTyrTyrLysTyrTyrMetLysLeuLysAsnPhe	280
Db	868	GTTGAGAGCTTACCGAAGCAGCAGCATGGTCTTCAAAAACCTGCACATCAATAATGAACCTTC	927
Oy	281	MetValAsnArgValPheIleProThrLysLysPheAsnLysGlnIleArgGluPro	300
Db	928	TTCGTGAACAGATTCATGAACCCACCAAGAGTTCTGTGAACAAGATCCATGACCC	987
Oy	301	SerLysAlaLeuLysGluLysValSerThrAspThrLysAspLeuPheGluAsnLysIle	320
Db	988	-----ACCAAGAGACTTCTTCGTGAACAAGATC	1011
Oy	321	GlyGlnGlyThrValAspPhePheAsnLysGluIleArgAspProSerLysAlaLeuLys	340
Db	1015	CATGAACCCACCAAGAGTTCTTCGTGAACAACATCCATGAACCCAC---	1067
Oy	341	GluLysValSerAsnAspAlaLysAspLeuPheGluAsnLysIleGlyGlnGlyThrVal	360
Db	1063	-----AAGCAATCTTGTGTGAACAAGTCCATGAACCCACCAAG	1107
Oy	361	AspPheIleAsnAsnGlnIleIleArgAspProSerLysAlaLeuIleArgLysValSerThr	380
Db	1102	GAGTTCTTCGTGAACAAGCTCCATGAACCCACCAAGAGTTCTTCCTAACAATGTGATACC	1161
Oy	381	GlyValIleGluAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPheIleAsnAsn	400
Db	1162	GGGGCATTTCCAGAAGATATCTGAAGAAGCAGTAGA-----	1197
Oy	401	GluIleArgAspProSerLysAlaLeuIleArgLysValIleThrGlnAlaAspLeu	420
Db	1198	CACCTGAAGAAGT---TCTAAGACAGTGTCCCGCAG-----GACAGGCCCTCATCGTCT	1244
Oy	421	PheGluAsnLysIle-----GlyGlnGlyThrVal---AspPheIleAsnLysGlu	436
Db	1249	CTGGAATAATGAGGCTGTAGAGATGGTCAACCTACATGAGGGAGATGTGACCATTTTGAA	1308
Oy	437	IleArgAspPro-----SerLysAlaLeuIleArgLysValSerThrGlu	451
Db	1309	ATGGCGACTCCCACTTGTAGCAGAGGCTCACAGAGAGATTTAATGAAAGTTGTGAACGA	1366

XX	Synthetic.
OS	
XX	
XX	repeat region; immune response; vaccine; ss.
KW	bbselosls; parasite; Babesia bovis; 21B4/rhoptry antigen; gene;
KM	Polymerase chain reaction; PCR; amplify; primer; detection;
DE	21B4/rhoptry antigen gene 5 DNA.
XX	
DT	13-JAN-1994 (first entry)
XX	
AC	AAO47075;
XX	
ID	AAO47075 standard; DNA; 1491 BP.
XX	
XX	
RESULT	7
AAO47075	

PN WD9314204 -A.  
XX  
PD 22-JUL-1993.  
XX  
PF 15-JAN-1993; 93WC-AU00012.  
XX  
PR 15-JAN-1992; 92AU-0000399.  
XX  
PA (CSIR ) COMMONWEALTH SCI & IND RES ORG.  
XX  
PI Dalrymple BP, Peters JW:  
DR WPI, 1993-243219/30.  
DR P-PADB; AAR39901.  
XX  
PT Detecting closely linked gene copies which encode protective  
PT antigen against babesiosis - by screening babesial genomic DNA  
PT library with oligo-nucleotide probe based partial sequencing of  
PT protective antigen and identifying positive clones  
XX  
SS Claim 23; Fig 6, 55pp: English.

CC This sequence represents the Babesia bovis 21B4/rhophy antigen gene  
CC 5. This sequence was isolated by PCR using the primers given in  
CC A0407068-72. Primer 21B4.1 corresponds to part of the repeated  
CC region of 21B4/rhophy antigen. In hybridisation assays this primer  
CC recognised two tandemly repeated regions suggesting that B. bovis  
CC contains two copies of the 21B4/rhophy antigen gene. The two  
CC proteins encoded by the two antigen genes are identical. Primers  
CC 21B4.2 and 21B4.3 flank the 21B4-309 coding region of the antigen  
CC gene. Primer 21B4.4 primes synthesis just 3' to the end of the open  
CC reading frame. The entire open reading frame was shown to encode five  
CC antigen genes. The 3 non-repetitive sequences of open reading frames  
CC 1-4 are identical. Gene 5 shows sequence divergence throughout most  
CC of the open reading frame. Babesia antigen genes can be used in the  
CC production of a combined vaccine which will stimulate a greater immune  
CC response and afford broader immunity than a single antigen vaccine.  
XX  
XX Sequence 1491 BP; 479 A; 286 C; 328 G; 398 T; 0 other;

Alignment Scores:	
Pred. No.:	3,45e-65
Score:	759.50
Percent Similarity:	54.04
Best Local Similarity:	34.268
Query Match:	32,208
DB:	14
Length:	149
Matches:	161
Conservative:	93
Mismatches:	173
Indels:	43
Gaps:	8

US-09-807-459-2 (1-458) x AAQ47075 (1-1491)

Oy	1	MetAlaIaP	roSerAspSerValGI	AspAlaThrI	hryThrLeuLeuAlaIaIaSerGI	User	20							
		91	CTAGACCTGCTGAAGTAGTGGTGATCTTACATATC	CACCTGAA	AAATTCGTGACGAAATT	150								
Db	21	ValAspSer	AlaIaAsnAlaTy	rMetIleasn	SerAspMetSerAspTy	rLeuSerAla	40							
		151	ATTAATGCCGAAACATRGAACATGAAATTAAT	TGCTGATATGCATTTA	AGACGTGTGAA	210								
Oy	41	ValSerAsp	AsnPhenAlaIu	ArgIle	oCSerGI	nAlaP	ProIyS	GI	AsnCyS	Ser	60			
		211	GAGGCACTAAATTTATTTGATCAATATTTGT	CAGGAAGTCGCTGAGAAATTC	TAAGTGGCT	270								
Db	61	AlaSerVal	SerAlaTy	rMetSerArgCys	AlaIyS	GI	nAspCyS	LeuThr	LeuGI	nSer	80			
		271	GACACAATTGAATCATATGTAAACGTTGTGA	GAGAAACAAC	TGTTTACAGATTG	ACGAA	330							
Oy	81	LeuIyS	TyrProLeu	GI	AlaIyS	TyrGI	nProLeu	ThrLeu	ProAsp	ProTy	rGI	nLeu	1000	
		331	GTAGCTTACCTTGAAATCAGCAATATCAGCA	CACTATTACTCCGGAACCAT	ACCAATTG	380								
Db	101	GI	AlaIaIaP	helleLeu	PhuIyS	GI	User	AspAla	nProAla	nSer	Thr	GI	Ulys	1200
		391	GATGTCGCTTACATATGTTTCACAAACGTG	AGCTATACCTGCTAA	AAAAATCGGCTGAA	450								



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OY 121 ArgPheTrpMetArgPheArgArgGlyLysAsnHisSerTyrPheHisAspLeuValPhe 140
DB 451 GGTCCATGGATGCGTACAGAGAGGATGAAGCATGGTGCATTCATCACA 510
OY 141 AsnLeuLeuGluLysAsnValThrArgAspAlaAspAlaThrAspIleGluAsnPheAla 160
DB 511 AGTTGGCTCGGCAGAGATGGTTCCTAAGATGCGTCTACTGACTTGATCCCGTC 570
OY 161 SerArgTyrLeuTyrMetLalaThrLeuTyrTyrLysThrTyrThrAsnValAspGluPhe 180
DB 571 AACAGCTTTGTACATGGCCACACCACTACTACAAAACCTACTAATGTTCAGAGAATTC 630
OY 181 GlyAlaSerPhePheAsnLysLeuSerPheThrThrGlyLeuPheGlyTrpGlyIleLys 200
DB 631 GGTGCTCGCTTTTCAACACTTCTCTTTACAGATATATATCGGTATGATTTAA 690
OY 201 ArgAlaLeuLysGluIleLeuArgSerAsnLeuProLeuAspIleGlyThrGluHisSer 220
DB 691 AGGGCATTTAAAGGATCGTCGCTCCATATGTTCCGGAAGACATGGGA---GAGCAGAT 747
OY 221 ValSerArgLeuGlnHisIleThrSerSerTyrLysAspTyrMetAspThrGlnIlePro 240
DB 748 ATTGAACGATATTAGTCAATTTGTCGGAAGGATACAGACTACATGTTCACACAGCTGCCA 807
OY 241 AlaLeuProLysPheAlaLysArgPheSerLeuMetValValGlnArgLeuLeuAlaThr 260
DB 808 ACCCTTTCAAGATTGCGCAACGCTTACTGACATGTTATGAACTTCTGTTGACACAG 867
OY 261 ValAlaGlyTyrValAspThrProTyrTyrLysLysTrpTyrMetLysLeuLysAsnPhe 280
DB 868 CTCGCCGGTTATGTCAAGCTCCATGCTACAGAGATGATGCAATTAATGATGCTCTTG 927
OY 281 MetValAsnArg-----ValPheIle 287
DB 928 TTAACGTGTGAAGCTTACCAACCTGATGAGATATACATTACTTAACTTATTTTCGTT 987
OY 288 ProThrLysLysPhePheAsnLysGluIleArgGluProSerLys---AlaLeuLysGlu 306
DB 988 GATACCTCCAGCAATTAACATAAAGATGACACTTAACCCAGCTGCTATGCTGAGAGAA 1047
OY 307 LysValSerThrAspThrLysAspLeuPheGluAsnLysIleGlyGlnGlyThrValAsp 326
DB 1048 AATATCGTTAACCCCGTCAGTATATCTCCGACGCCAACAATATTTCTAGTGCACA 1107
OY 327 PhePheAsn-----LysGluIleArgAspProSerLysAlaLeu 339
DB 1108 AACTACAAATGACGGCATATAAATAGATCCCTCTTTATGAACTTAAGAGCGGCAT 1167
OY 340 LysGluLysValSerAsnAspAlaLysAspLeuPheGluAsnLysIleGlyGlnGlyThr 359
DB 1168 ATCGGAATTGCTGCGATACCGCTAGCATTAATATAGATATAGATAAGTAATAA----- 1221
OY 360 ValAspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLysValSer 379
DB 1222 -----GCGAAAGAATTAAGTGCCTGCCAAG 1248
OY 380 ThrGlyAlaGluAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPheIleAsn 399
DB 1249 GACCGCGCACAGGCAATATATGCAAGATCAAGTGAACCTGCTGTAGTATATTAACGAC 1308
OY 400 -----AsnGluIleArgAspProSerLysAlaLeuIleArgLysValTyr----- 414
DB 1309 GTTGTAAAGATATGATCTCTTGATGCAGTAAC-----ATCAGAAATATATTAACGTGC 1362
OY 415 ---ThrGluAlaAspAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPheIle 433
DB 1363 AGCTTCAAGATGACAAATATATGAAACAAGACTGAGAGAGCAAAAGTTGAGGAAGTT 1422
OY 434 AsnLysGluIleArgAspProSerLysAla 443
DB 1423 AAACCTGAGCTGAAGCAAAAAGATGTGCT 1452

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RESULT 8
AAQ26065
ID AAQ26065 standard; DNA; 282 BP.
XX
AC AAQ26065;
XX
DE 09-DEC-1992 (first entry)
XX
DE 21B4 gene clone pT#13, EcoRI insert.
XX
KW Beta-galactosidase; B. bovis; Bb; T21B4; ss.
XX
OS Babesia bovis.
XX
PN EP492525-A.
XX
PD 01-JUL-1992.
XX
PF 20-DEC-1991; 91BP-0121990.
XX
PR 21-DEC-1990; 90AU-0004051.
XX
PA (CSIR ) COMMONWEALTH SCI & IND RES ORG.
XX
PI Casu RE, Commins MA;
XX
DR WPI, 1992-218727/27.
XX
DR P-PSDB; AAR25187.
XX
PT Monoclonal antibody to Babesia bovis parasite - used to isolate
PT antigens for use in vaccines for treating Babesiosis and
PT providing immunity in cattle
XX
PS Claim 13; Fig 9; 24pp; English.
XX
CC The sequences given in AAQ26062 and AAQ26065-7 are portions of the 21B4
CC gene which were isolated from a B. bovis (Bb) cDNA lambda gt11
CC library and cloned into pGEM7zf(+). The resulting plasmids were
CC transformed into E. coli strain JM83. The inserts were in frame,
CC when translated, with the vector beta-galactosidase gene. The fusion
CC proteins produced by translation of these vectors were recognised by
CC the monoclonal antibody of the invention, T21B4. These fusion
CC antigens could be used in vaccines for the treatment of babesiosis
CC and to provide immunity in relation to Bb infection in cattle against
CC different strains of Babesia by heterologous and homologous challenge.
XX
SQ Sequence 282 BP; 80 A; 60 C; 63 G; 79 T; 0 other;

Alignment Scores:
Pred. No.: 3.18e-07 Length: 282
Score: 161.00 Matches: 32
Percent Similarity: 51.65% Conservative: 15
Best Local Similarity: 35.16% Mismatches: 44
Query Match: 6.82% Indels: 0
Gaps: 0

US-09-807-459-2 (1-458) x AAQ26065 (1-282)
OY 2 AlaProSerAspSerValGlyAspValThrLysThrLeuLeuAlaIleSerGluSerVal 21
DB 10 GCTCCAGCTGAAAGTGAAGTGAATTAATCTCCACATTTGGAACAGCTGATTTGATG 69
OY 22 AspSerAlaIleAsnAlaTyrMetIleAsnSerAspMetSerAspTyrIleSerAlaVal 41
DB 70 ACTTCTCGTGACCAACATGCAACACATTACTAAGGATATGAAACAGCTTTGACCAATGCT 129
OY 42 SerAspAsnPheAlaGluArgIleCysSerGlnValProLysGlySerAsnCysSerAla 61
DB 130 CGTGACAGATGTGAATATGATGTTTGCTTAATGCTCTGAGACATCCCACTGCGTAG 189
OY 62 SerValSerAlaTyrMetSerArgCysAlaLysGlnAspCysLeuThrLeuGlnSerLeu 81
DB 190 GTAGTTAAACAACATATGCTGACCGTGTGAATATGACGATCTTACAGATTGACAAATGTC 249

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OY      82 LysTyrProLeuGluValIleuTyrGlnProLeu 92
        :::::::::::::::::::::
DB      250 AGATATCCCTGTACCAAGATACCAACTCTTA 282

RESULT 9
AAV21209
ID      AAV21209 standard; DNA; 1664976 BP.
AC      AAV21209;
DT      10-NOV-1998 (first entry)
DE      Methanococcus jannaschii circular chromosome.
XX
XX      Methanococcus jannaschii; methanogenic archaeon; circular chromosome;
KM      genome; autotrophic; extrachromosomal element; identification; ds.
XX
OS      Methanococcus jannaschii.
XX
XX      WO9807830-A2.
XX
XX      26-FEB-1998.
XX
XX      22-AUG-1997; 97WO-US14900.
XX
XX      22-AUG-1996; 96US-0024428.
XX
XX      (GENO-) INST GENOMIC RES.
PA      (UNIT ) UNIV ILLINOIS FOUND.
PA      (UYXO ) UNIV JOHNS HOPKINS SCHOOL MEDICINE.
XX
XX      Bult CJ, Smith HO, Venter JC, White OR, Woese CR;
DR      WPT: 1998-169145/15.
XX
XX      Complete genome sequence of methano-genic archaeon, Methanococcus
PT      jannaschii - useful in identification of M. jannaschii genome
PR      fragment
XX
XX      Claim 13; Page 152-585; 614pp; English.
XX
XX      The present sequence represents the complete 1.66-megabase pair genome
CC      sequence of the Methanococcus jannaschii circular chromosome. The
CC      present invention describes M. jannaschii open reading frames from the
CC      genome sequence. The invention also describes a computer based system
CC      for identifying fragments of the M. jannaschii genome that are
CC      homologous to target nucleotide sequences, comprising: (a) data storage
CC      means comprising the nucleotide sequence of the 1664976, 58407 or 16550
CC      bp sequence (see AAV21209, AAV21210 and AAV21211), or a nucleotide
CC      sequence at least 99.9% identical to it; (b) search means for comparing a
CC      target sequence to the nucleotide sequence of the data storage means to
CC      identify a homologous sequence, and (c) retrieval means for obtaining
CC      the homologous sequence. The method, which is based on whole genome
CC      random sequencing of an autotrophic archaean M. jannaschii, the genome
CC      of which consists of 3 physically distinct elements, a large circular
CC      chromosome (the 1664976 bp sequence given in AAV21209), a large circular
CC      extra-chromosomal element (the 58407 bp sequence given in AAV21210), and
CC      a small circular extra-chromosomal element (the 16550 bp sequence given
CC      in AAV21211), can be used in the identification of M. jannaschii genome
CC      fragment.
XX
XX      Sequence 1664976 BP; 568133 A; 264649 C; 258701 G; 573392 T; 101 other;
SQ

Alignment Scores:
Pred. No.: 0.823 Length: 1664976
Score: 147.00 Matches: 88
Percent Similarity: 38.79% Conservative: 59
Best Local Similarity: 23.22% Mismatches: 150
Query Match: 6.23% Indels: 82
DB: 19 Gaps: 17

US-09-807-459-2 (1-458) x AAV21209 (1-1664976)

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Oy 440 roSerLyAlaLeuIleArgLyValSerThrGluAlaAspAsnLeuGlu 457
Db 1419240 TTGGGGGAGAAATTGTTAAATAATACATTAAGTGTAGAG---TTGTTAGAG 1419289
RESULT 10
ID AAQ26066 standard; DNA: 170 BP.
XX
AC AAQ26066:
XX
DT 09-DEC-1992 (first entry)
XX
DE 21B4 gene clone PT#13, EcoRI insert (2).
XX
KM Beta-galactosidase; B. bovis; Bb; T21B4; ss.
XX
OS Babesia bovis.
XX
PN EP492525-A.
XX
PD 01-JUL-1992.
XX
PF 20-DEC-1991; 91EP-0121990.
XX
PR 21-DEC-1990; 90AU-0004051.
XX
PA (CSIR ) COMMONWEALTH SCI & IND RES ORG.
XX
PI Casu RE, Commings MA;
XX
DR WPI: 1992-218727/27.
XX
P-PSDB: AAR25188.
XX
PT Monoclonal antibody to Babesia bovis parasite - used to isolate
PT antigens for use in vaccines for treating Babesiosis and
PT providing immunity in cattle
XX
PS Claim 15; Fig 10; 24pp; English.
XX
CC The sequences given in AAQ26062 and AAQ26065-7 are portions of the 21B4
CC gene which were isolated from a B. bovis (Bb) cDNA lambda gt11
CC library and cloned into pGEM7zf(+). The resulting plasmids were
CC transformed into E. coli strain JM83. The inserts were in frame,
CC when translated, with the vector beta-galactosidase gene. The fusion
CC proteins produced by translation of these vectors were recognised by
CC the monoclonal antibody of the invention, T21B4. These fusion
CC antigens could be used in vaccines for the treatment of babesiosis
CC and to provide immunity in relation to Bb infection in cattle against
CC different strains of Babesia by heterologous and homologous challenge.
XX
SQ Sequence 170 BP; 56 A; 28 C; 31 G; 51 T; 4 other:
XX
Alignment Scores:
Pred. No.: 2.98e-05 Length: 170
Score: 138.00 Matches: 27
Percent Similarity: 66.67% Conservative: 9
Best Local Similarity: 50.00% Mismatches: 18
Query Match: 5.838 Indels: 0
Gaps: 0
DB: 13
US-09-807-459-2 (1-458) x AAQ26066 (1-170)
Oy 161 SerAlaGlyrLeuTyrMetAlaThrLeuTyrTyrThrAsnValAspGluPhe 180
Db 1 AACAAAGTACTTAAATGCTACATGACCTCAAGACTTATTGACACTAAACAGTANG 60
Oy 181 GlyAlaSerPheAsnLysLeuSerPheThrThrGlyLeuPheGlyTrrpGlyIleLys 200
Db 61 AACGCCAANTTCTTCAACAGATTCAGCTTTACTACAAAGATTTTCAGTNNTCGTATTAGG 120
Oy 201 ArgAlaLeuLysGlnIleLeuArgSerAsnLeuProLeuAsp 214
Db 121 CAATCATTTGAGTATATCATCTCAGGTGGAATGTCTTCTGAAGAT 162

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RESULT 11
ID AAX20254/c standard; DNA: 26776 BP.
XX
AC AAX20254;
XX
DT 04-MAY-1999 (first entry)
XX
DE Borrelia burgdorferi polynucleotide sequence #7.
XX
KM Borrelia burgdorferi; spirochete; bacterium; pathogen; Lyme disease;
KM epidemic relapsing fever; endemic relapsing fever; Lyme borreliosis;
KM infection; diagnosis; characterisation; detection; ds.
XX
OS Borrelia burgdorferi.
XX
PN W09858943-A1.
XX
PD 30-DEC-1998.
XX
PF 18-JUN-1998; 98WO-US12764.
XX
PR 03-SEP-1997; 97US-0057483.
PR 20-JUN-1997; 97US-0050359.
PR 22-JUL-1997; 97US-0053344.
PR 22-JUL-1997; 97US-0053377.
XX
PA (HUMA-) HUMAN GENOME SCI INC.
PA (MEDI-) MEDIMUNE INC.
XX
PI Clayton R, Dougherty BA, Fraser C, Lathigra R, Smith HO;
PI White OR;
XX
DR WPI: 1999-081217/07.
XX
PT New isolated Borrelia burgdorferi nucleic acids - used to develop
PT products for the detection, diagnosis, characterisation, prevention
PT and therapy of infections, particularly Lyme disease
XX
PS Claim 1; Page 867-882; 1128pp; English.
XX
XX
AAAX20248 to AAX20402 represent polynucleotide sequences isolated from
CC Borrelia burgdorferi (Bb). Products derived from Bb can be used for
CC the detection, diagnosis, characterisation, prevention and therapy of
CC Bb infections, e.g. Lyme disease. They can also be used for the
CC production of biosynthetic products, e.g. enzymes. Borrelia belongs
CC to a family of motile, spiral-shaped bacteria called Spirochetes.
CC Spirochetes are pathogenic in humans and Borrelia causes epidemic and
CC endemic relapsing fever, and Lyme borreliosis, more commonly known as
CC Lyme disease.
XX
SQ Sequence 26776 BP; 8604 A; 4331 C; 4257 G; 9580 T; 4 other:
XX
Alignment Scores:
Pred. No.: 0.0633 Length: 26776
Score: 134.00 Matches: 61
Percent Similarity: 40.96% Conservative: 75
Best Local Similarity: 18.37% Mismatches: 124
Query Match: 5.68% Indels: 72
Gaps: 13
DB: 20
US-09-807-459-2 (1-458) x AAX20254 (1-26776)
Oy 167 AlaThrLeuTyrTyrLeuThrThrThrAsnValAspGluPheGlyAlaSerPheAsn 186
Db 6709 GCTACAAATTAATCTTAACATC-----AAATTTGGGTGAG 6677
Oy 187 LysLeuSerPheThrThrThrLeuPheGlyTrrpGlyIleLysArgAlaLeuLysGlnIle 206
Db 6676 CTGTAGCATATCTTGCCCGCAAGTGGTGAAGGCAATGTTGTTCAAGGCCAATT 6617
Oy 207 IleArgSerAsnLeuProLeuAspIleGlyThrGlnHisSerValSerArgLeuGlnHis 226

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Db 6616 ACCAGCCCTTTGTTAAAGCTCTACGAGGAAATACGTAATCGGCACACCTAAAGAA 6557
Oy 227 TLehrserSerTyrlsAspThrMetAspThrGlnIleProAlaLeuProLysPheAla 246
Db 6556 CTTCGAACA-----TATGTTCTTAGTGATTT-----TTAGATCGTTTAAAG 6515
Oy 247 LysArgPheSerLeuMetVal-----ValGlnArgLeuAlaThrValAlaGlyTyr 264
Db 6514 GGTCGGTTTGAGCCATTATCAAAATGGTTCAAAAGTTGATGATGATATCC----- 6461
Oy 265 ValAspThrProTryptylsLysTryptylMetLysLeuLysAsnPheMetVal----- 282
Db 6460 -----AAAGCGTATACAAATTTGAAATTCGTGTTACTTTTGG 6422
Oy 283 -----AsnArgValPhe 286
Db 6421 AAAAAGCCGAAAGAAAGAAAGACACTTCTGCTCCAGACCCGAAAGGATTAAGTTT 6362
Oy 287 TLeProThrLysLysPhe---PheAsnLysGlnIleArgLysProSerLysAlaLeuLys 305
Db 6361 GATCCAAATGCAAAACCAATATTAACAGAAATGGCAGAAACCTATCAAAACCTCGAA 6302
Oy 306 GlnLysValSerThrAspThrLysAspLeuPheGlnLysLysIleGlyGlnGly----- 323
Db 6301 GACGAAATATTTAATAGCGCAGAGGACATCTAT--AATAAGACCGGAAAGCAGAGAA 6245
Oy 324 -----ThrValAspPhePheAsnLysGlnIleArgAspPro 335
Db 6244 CAAGCCTTAAGATCTTGAAAAAACCATTAATGAAAAAACCAAAAGTTTAAGATGAA 6185
Oy 336 SerLysAlaLeuLysGlnLysValSerAsnAspAlaLysAspLeuPhe----- 351
Db 6184 TACTCCAAATATTTGATTCAGTTAAGTACGAGCAACAAATTTAGTCGAGATTGAA 6125
Oy 352 -----GlnLysLysIleGlyGlnGlyThrValAspPheIleAsnGlnIleArgAsp 369
Db 6124 AATCACTAATATGATTCATTAACCTCAATATGATTTTGGAATGAG----- 6077
Oy 370 ProSerLysAlaLeuIleArgLysValSerThrGlnLysAlaGlnAspLeuPheGlnLys 389
Db 6076 ---TACCAAAATTTACTTAAAGAAAGAAAGAGTGTGAAGGAAATATCAAAACTCG 6020
Oy 390 TLeGlyGlnGlyThrValAspPhe-----IleAsnAsnGlnIleArgAspProSer 406
Db 6019 CCCCATACAGATCAAGTACGCTTTACAGAAACCTTAATGATGAAATCAAGAAAGAAC 5960
Oy 407 LysAlaLeuIleArgLysValTyrThrGlnAlaAspAspLeuPheGlnLysIleGly 426
Db 5959 AAAGCGTTTCGTAAGAAATATGAAAAAGTTTCGAACCTCG-----AACGAGCTTAAC 5906
Oy 427 GlnGlyThrValAspPheIleAsnLysGlnIleArgAspProSerLysAlaLeuIleArg 446
Db 5905 AGGCAGATTGTAGTCGACCTTGAAGAACAGGTTATGATGATGAAAAACCGCTTTGAT 5846
Oy 447 LysValSerThrGlnAlaAspAsnLeuLeuLys 458
Db 5845 CGATCTTTTGTGAGGCTCAAAAGCTCTGCAAAA 5810

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RESULT 12  
AAx20250/c  
ID AAx20250 standard; DNA: 111309 BP.

XX AC AAx20250;  
XX DT 04-MAY-1999 (first entry)

DE Borrelia burgdorferi polynucleotide sequence #3.

XX KW Borrelia burgdorferi; spirochete; bacterium; pathogen; Lyme disease;  
XX KM epidemic relapsing fever; endemic relapsing fever; Lyme borreliosis;  
XX infection; diagnosis; characterisation; detection; ds.

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OS Borrelia burgdorferi.
FN
XX MO9858943-A1.
XX
XX 30-DEC-1998.
XX
XX 18-JUN-1998; 98WO-0512764.
XX
XX 03-SEP-1997; 97US-0057483.
XX 20-JUN-1997; 97US-0050359.
XX 22-JUL-1997; 97US-0053344.
XX 22-JUL-1997; 97US-0053377.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX (MEDI-) MEDIMUNE INC.
XX
XX Clayton R, Dougherty BA, Fraser C, Lethigra R, Smith HO:
XX white OR;
XX
XX WPI; 1999-081217/07.
XX
XX New isolated Borrelia burgdorferi nucleic acids - used to develop
XX products for the detection, diagnosis, characterisation, prevention
XX and therapy of infections, particularly Lyme disease
XX
XX Claim 1; Page 738-800; 1128pp; English.
XX
XX AAx20248 to AAx20402 represent polynucleotide sequences isolated from
XX CC Borrelia burgdorferi (Bb). Products derived from Bb can be used for
XX CC the detection, diagnosis, characterisation, prevention and therapy of
XX CC Bb infections, e.g. Lyme disease. They can also be used for the
XX CC production of biosynthetic products, e.g. enzymes. Borrelia belongs
XX CC to a family of motile, spiral-shaped bacteria called Spirochetes.
XX CC Spirochetes are pathogenic in humans and Borrelia causes epidemic and
XX CC endemic relapsing fever, and Lyme borreliosis, more commonly known as
XX CC Lyme disease.
XX
XX SQ Sequence 111309 BP; 35956 A; 13151 C; 19075 G; 43117 T; 10 other:
XX
XX
XX Alignment Scores:
XX Pred. No.: 5.75 Length: 111309
XX Score: 122.50 Matches: 93
XX Percent Similarity: 37.47% Conservative: 55
XX Best Local Similarity: 23.54% Mismatches: 144
XX Query Match: 5.19% Indels: 103
XX DB: Caps: 20
XX
XX US-09-807-459-2 (1-458) x AAx20250 (1-111309)
XX
XX Oy 113 AsnProAlaAsnSerThrGlnLysArgPheThrMetArgPheArgGlyLysAsnHis 132
XX ||||| |||
Db 47193 AACCCAAATTAACCAAGAAATTAACACATTT----- 47164
Oy 133 SerTyrlsAspLeuValPheAsnLeuLeuGlnLysAsnValThrArgAspAlaAsp 152
XX ||||| |||
Db 47163 AAACCTTTTAACTTATGATTTTATATATCTTAAAGAAATTTTAAACAGAGAA----- 47110
Oy 153 AlaThrAspIleGlnAsnPheAlaSerArgTyrLeuTyrMetAlaThrLeuTyrTyrLys 172
XX ||||| |||||
Db 47109 -----GCCTAGAAATATTTTAAAGAAAGCTTTTCTGATTAATAAA 47071
Oy 173 ThrTyrThrAsnValAspGluPhe---GlyAlaSerPhePheAsnLysLeuSerPheThr 191
XX ||| : : : : : |||
Db 47070 ACAGATGAAGATGCMAATTAATATCTTGTGTACATTTTAAAGAAAGATGATATAAA 47011
Oy 192 ThrGlyLeuPheGlyTyrPglyTleLysArgAlaLeuLysGlnIle----- 206
XX ||| : : : : : |||||
Db 47010 GAACCTCTATAC-----TATCTTAAAAAGTCCCAATTTTAATAAA 46969
Oy 207 -----IleArgSerAsnLeuProLeuAspIleGlyThrGlnHisSerValSer 222
XX : : : : : |||
Db 46968 AATAATAAGATGTCGTAATAAACACCTGGAAATTAACCTTATTAATCTGAAGAACTACGA 46909

```



QY	263	GLYThrValAspThrProTrpTyrLysLysTrpTyrMetLysLeuLysAsnPhenMetVal	282
Db	636234	TTGGCAATAGCTTAATCTTTATATCAAAATAATCAAGATCATCTACGACTAGAGATTGCC	636175
QY	263	GLYThrValAspThrProTrpTyrLysLysTrpTyrMetLysLeuLysAsnPhenMetVal	282
Db	636174	AACCAAAATA	636145
QY	283	AsnArgValPheIleProThrLysLysPhePhe-----AsnLysGluIleArgGluPro	300
Db	636144	TATGAGCGCTTTATGATTACATCTGCAATTCATTCAAAAAATTAAGAAATTAGAAAAATTA	636085
QY	301	SerLysAlaLeuLysGlu-----LysValSerThrAspThrLys-----	313
Db	636084	GAACAAATATTTAAAGAAATATATAAAGTAAACCACGACTTGCTTAATAAACTTTCCT	636025
QY	314	AspLeuPheGluAsnLysIleGlyGlu-----	323
Db	636024	GAATATTATTATAT-----CTTGAGAACTCAATCTCTGTGAAAAATTATCAAAAGCT	635968
QY	324	ThrValAspPhePheAsnLysGluIleArgAspProSer-----LysAlaLeuLysGluLys	342
Db	635967	ACAGAGACTTTTACCAAGTTGAAGAGATTGATTCCAATATTAATAAATATTAAGAAAA	635908
QY	343	ValSerAsnAspAlaLysAspLeuPheGluAsn-----LysIle	355
Db	635907	TTG---GAATTTACCAAAAGCTTTAAACGAAACATACCTTAAGATATATTTGGCCAGT	635851
QY	356	GlyGlnGlyThrValAspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIle	375
Db	635850	TCAAAAGCAAAATTTTGAATAATGCAAAATGAAATTT-----ATT	635812
QY	376	ArgLysValSerThrGlyAlaGluAspLeuPheGluAsnLysIleGlyGlnGlyThrVal	395
Db	635811	TTTAAACATATTTTAAATTAATTTCAATATAGAGATTCTAAATTAATGAATGAATACACA	635752
QY	396	AspPheIle-----AsnAsnGluIleArgAspProSerLysAla	408
Db	635751	CAATTTATATGACAAACTTTCACCTTAGCTACATACATACAGTGGGAGAGA-----AATTTA	635698
QY	409	LeuIleArgLysValTyrThrGluAlaAsp-----AspLeuPheGluAsnLysIle	425
Db	635697	ATAGTGCCTTTTGAACAGCAAGATACATCTTTGGTCAATTAATTC-----	635650
QY	426	GlyGlnGlyThrValAspPheIleAsnLysGluIleArgAspProSerLysAlaLeuIle	445
Db	635649	-----TTAAAGATTTTCATTTCATTTCAAAAATAAAGCAAAATTAATAAAGACATCTGT	635599
QY	446	-----ArgLysValSerThrGluAlaAspAsnLeuLeuGlu	457
Db	635598	ATTGCTCCATCAAAATCTCTTTAAAGGTCAAAACAAATGATTGA	635554
RESULT 14			
ID	AAAI5186	standard; DNA: 1165 BP.	
AC	AAAI5186;		
XX	04-SEP-2000	(first entry)	
DE	DNA encoding	Escherichia coli	virulence proteins.
XX	Virulence protein:	tatA; tatB; tatC; tatE; mdgG; creG; yggN;	
KW	ecoli; ltrD; ltrC; ltrE; mtd2; msl; vaccine; infection;		
KW	Grim negative bacterium; ss.		
OS	Escherichia coli.		
XX	key	location/Qualifiers	
FT	FT	2..1099	
CT	CT	/*tag= a	
FT	FT	/product= "virulence protein"	

FT		/note= "encodes AAY93237; no termination codon given"
FT	CDS	1102..1488
FT		/*tag= b
FT		/product= "virulence protein"
FT		/note= "encodes AAY93238"
FT	CDS	1573..1896
FT		/*tag= c
FT		/product= "virulence protein"
FT		/note= "encodes AAY93239"
FT	CDS	1939..2196
FT		/*tag= d
FT		/product= "virulence protein"
FT		/note= "encodes AAY93240; no termination codon given"
FT	CDS	2198..2533
FT		/*tag= e
FT		/product= "virulence protein"
FT		/note= "encodes AAY93241"
FT	CDS	2613..3041
FT		/*tag= f
FT		/product= "virulence protein"
FT		/note= "encodes AAY93242; no termination codon given"
FT	CDS	3054..3410
FT		/*tag= g
FT		/product= "virulence protein"
FT		/note= "encodes AAY93243"
FT	CDS	3460..3705
FT		/*tag= h
FT		/product= "virulence protein"
FT		/note= "encodes AAY93244"
FT	CDS	3791..4837
FT		/*tag= i
FT		/product= "virulence protein"
FT		/note= "encodes AAY93245"
FT	CDS	4878..7802
FT		/*tag= j
FT		/product= "virulence protein"
FT		/note= "encodes AAY93246"
FT	CDS	7816..9483
FT		/*tag= k
FT		/product= "virulence protein"
FT		/note= "encodes AAY93247"
FT	CDS	9836..10084
FT		/*tag= l
FT		/product= "virulence protein"
FT		/note= "encodes AAY93248"
FT	CDS	10134..10430
FT		/*tag= m
FT		/product= "virulence protein"
FT		/note= "encodes AAY93249"
FT	CDS	10459..10779
FT		/*tag= n
FT		/product= "virulence protein"
FT		/note= "encodes AAY93250"
XX		
PN	WO200028038-A2.	
PD	18-MAY-2000.	
XX		
PF	09-NOV-1999;	99MO-GB03721.
XX		
PR	09-NOV-1998;	98GB-0024569.
PR	09-NOV-1998;	98GB-0024570.
PR	17-DEC-1998;	98GB-0027814.
PR	17-DEC-1998;	98GB-0027815.
PR	17-DEC-1998;	98GB-0027816.
PR	17-DEC-1998;	98GB-0027818.
PR	13-JAN-1999;	99GB-0000708.
PR	13-JAN-1999;	99GB-0000710.
PR	13-JAN-1999;	99GB-0000711.
PR	28-JAN-1999;	99GB-0001915.
PA	(MICR-) MICROSCIENCE LTD.	
XX		

PI Crooke RH, Clarke EE, Everest PH, Dougan G, Holden DM, Shea JE,  
PI Feldman RG,  
XX  
XX WPI: 2000-376550/32.  
DR P-PSDB: AAY93237, AAY93238, AAY93239, AAY93240, AAY93241, AAY93242,  
DR AAY93243.  
XX  
XX Peptide encoded by an operon including genes from *Escherichia coli* for  
PT screening potential drugs, detecting virulence and treating conditions  
PT associated with infection by a Gram negative bacterium -  
XX  
PS Disclosure: Page 83-101; 122pp: English.  
XX  
XX The present sequence encodes *Escherichia coli* virulence proteins  
CC The specification describes virulence proteins which are encoded  
CC by an operon including *tatA*, *tatB*, *tatC*, *tatE*, *mdcB*, *creC*, *recG*, *y99N*,  
CC *ecfI*, *iroD*, *iroC*, *iroB*, *mtd2* or *msl-16* genes obtained from *Escherichia*  
CC *coli* K1. The virulence proteins and polynucleotides, and their vaccines  
CC are useful for screening potential drugs, for the detection of virulence  
CC and for treating or preventing conditions associated with infection by  
CC a Gram negative bacterium particularly *Escherichia coli*.

OY	254	-----ValGlnATgLeuLeuAlaThrVal-----	261
Db	6093	TTTTTTCATATCGACGCTTGTGGAGCATGAAATTAAACAAAGAAACAATGCTTCA	6155
OY	262	-----AlaGlyTyrValAspThrProTyrIleLysIleStrPtyr	274
Db	6153	TTAGTGTAAAGAGCGTGCGGAGAAAGATTCATTGAAAATTAAGCCATATTAACAGTTTAA	6212
OY	275	MethylSerLeuLysAsnPhenMetVal-----	282
Db	6213	GAATATGCTTGAATAATTTCTTCATCGCAAAATCGCAAGAGATTCTTTAAGACGATGGCGAT	6272
OY	283	---AsnArgValIlePheIleProThrLysLysPheAsnLysGlu-----IleArg	298
Db	6273	ATGTCATGATTAATTGTTGTGACAAACCAACGATGAGAAATGAAACATCATGTTTATCT	6333
OY	299	GluProSerLysAlaLeuLysGluLysValSerThrAspThrLysAspLeuPheGluAsn	318
Db	6333	AAAAGATCTAAACAACTAAAGCTAAAGAAAAAGATAGATTAAAGAAAGATCTGTAT-----	6388
OY	319	LysIleGlyGlnGlyThrValAspPheAsnLysGluIleArgAspProSerLysAla	338
Db	6387	-----GATTTTTTGATAAGATTAGATATGATTACTGGAATATT	6425
OY	339	LeuLysGluLysValSerAsnAspAlaLysAspLeuPheGluAsn---LysIleGlyGln	357
Db	6426	GAATAAATAAGTCAATCAATAAACGAGAGATATTTCTCCAGTACACAAATTAACACAG	6485
OY	358	GlyThrValAspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLys	377
Db	6486	ACCATATATAGATTATGATATCAATAAATTAAGAACAAATGATCTCATTTAAATAT	6544
OY	378	ValSerThrGlyAlaGluAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPhe	397
Db	6546	CTACGTAATTCCTGTGGAT-----ATAAAGAAACCTCTGGAGCTGATTAACAAAGAG	6599
OY	398	IleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLysValIleThrGluAla	417
Db	6600	TTATCTAT---TTATGGATAGATATCAATTAAGAAACAAATAACTGTTATTCACATA	6655
OY	418	AspAspLeuPheGluAsn-----LysIleGlyGlnGlyThrValAsp	431
Db	6657	AATGAGCTAAAGATACGTTGATAGAAAGCTATAGAACTGATATTAATAATATGAT	6710
OY	432	PheIleAsn-----LysGluIleArgAsp-----	439
Db	6717	TTTCCACTTACGGAAGAGACTGAAGATCTTTGAACTACACMAAGTTACTATGAA	6776
OY	440	-----ProSerLysAlaLeuIleArgLysValSerThrGlu	451
Db	6777	AAAGACTTAACAAAGTTATATATAGACGCTAAATAATGCTTTGAAACATGTGCATCTTAA	6835
OY	452	AlaAspAsnLeuLeu	456
Db	6837	GCAAAATAGCTTAATT	6851
RESULT 15			
AAV71915			
ID	AAV71915 standard; DNA; 3883 BP.		
XX	AAV71915;		
XX	11-FEB-1999 (first entry)		
DE	S. cerevisiae CIN8 DNA sequence.		
XX	TIH1; recombinant; research; epitope mapping; modulating; CKI; yeast;		
KW	casein kinase I; cell growth; CIN8; cancer; viral infection; ss.		
OS	Saccharomyces cerevisiae.		
PN	US5846764-A.		



XX 08-DEC-1998.  
 PD 23-JAN-1995: 95US-0376843.  
 XX PF 23-JAN-1995: 95US-0376843.  
 XX PR 21-JAN-1994: 94US-0184605.  
 XX PA (ICOS-) ICOS CORP.  
 XX PI Demaglo AJ, Hoekstra ME;  
 XX DR WPI; 1999-059057/05.  
 XX PT Polynucleotide encoding yeast T1H1 polypeptide - useful for  
 XX PT producing recombinant polypeptide  
 XX PS Example 3; Columns 239-244 (columns 65-70); 46pp; English.  
 CC This represents the DNA sequence of *Saccharomyces cerevisiae* C1N8 that  
 CC belongs to the yeast kinesin gene family. The invention relates to T1H  
 CC proteins that interact with casein kinase I (CKI) isoforms. A host cell  
 CC transformed with a DNA expression construct containing the T1H DNA can  
 CC be used to produce recombinant T1H1 polypeptide which is used for  
 CC research purposes. The T1H proteins are used to map T1H epitopes in the  
 CC development of epitope-specific agents that may be useful for modulating  
 CC CKI/T1H protein interactions involved in cell growth in health and  
 CC disease, e.g. cancer or viral infections.  
 CC Note: The specification has erroneous column number indications.  
 CC  
 SO Sequence 3883 BP; 1495 A; 652 C; 680 G; 1056 T; 0 other:  
 Alignment Scores:  
 Pred. No.: 0.115 Length: 3883  
 Score: 120.00 Matches: 121  
 Percent Similarity: 34.75% Conservative: 92  
 Best Local Similarity: 19.74% Mismatches: 214  
 Query Match: 5.09% Indels: 186  
 DB: Gaps: 29  
 US-09-807-459-2 (1-458) x AAV71915 (1-3883)  
 QY 4 SerAspSerValGlyAspValThrLysThrLeuLeuAlaIleSerGluSerValAspSer 23  
 DB 1051 AGCGATTCACAGCAAAATTAACGAGGGTCTTTGAAGTTGTTGACACATTGGAATA 1110  
 QY 24 AlaAlaSerAlaLysMetIle-----AsnSerAspMetSer 35  
 DB 1111 CAACAGACCAATTACGATAGTAATAATGTCCTTCACTTCACTCAACGAAATTAAG 1170  
 QY 36 AspTyrLeuSerAlaValSerAsp-----AsnPheAla 46  
 DB 1171 GACCTCTTGACAGCAATACCAACGGCTCTAGTAATACGTGGCTTGAAGGCAATTAATG 1230  
 QY 47 GluArgLleCysSerGlnValProLysGlySerAsnGlySerAlaSerValSerAlaTyr 66  
 DB 1231 AAAAATTTGAGATTTTGGCTTCACAGCAACAAATATACCTACCAACAGAGTCTAGT 1290  
 QY 67 MetSerArgCysAlaLysGlnAspCys-----LeuThrLeuGln 79  
 DB 1291 AGTTCCAGAGTAATTTCTAGGACAGTTCTCCAGGTCATTAATGATCTPAACACCTPAA 1350  
 QY 80 Ser-----LeuLysTyrProLeuGluAlaLysTyrGlnProLeuThrLeuProAspPro 97  
 DB 1351 GCGTCTCTAATTAAGAAAAGTTAAGACAAATATCACTGCGCATACATCAACACACAG 1410  
 QY 98 TyrGlnLeuGluAlaIlePheIleLeuPheLysGluSerAspAlaAsnProAlaAsnSer 117  
 DB 1411 TATCAACCAACAGCAGCAGTAATTCAGAGCAACACTCTCTACTGCTGCTTACC 1470  
 QY 118 ThrGlu-----LysArgPheThrPmeArgPhe 126  
 DB 1471 ACTAATATATGCTTCTACTAACCACCAACCAAAATTAACGCTCAAGAAGTTCAGTGTCCA 1530

QY 127 ArgArgGlyLysAsnHisSerTyrPheHisAspLeuVal----- 139  
 DB 1531 AATGACCAAACTATATGCTATATACATCCAGAAATTTGCAGAAATTTACATAAACATGCT 1590  
 QY 140 -----PheAsnLeuLeuGluLysAsnValThr---ArgAspAlaAspAlaThrAsp 155  
 DB 1591 ATGGAGGGGCTAAACCTATTACAAAAGGCTTAAAGCATAGGCAAGTAGGCTCCATAAA 1650  
 QY 156 IleGluAsnPheAlaSerArg-----TyrLeuTyrMetAlaThrLeuTyrTyrLysThr 173  
 DB 1651 ATGACGATTTTTCACATAGATTCATACATTTTTCACATCTTGTATAGAACAT 1710  
 QY 174 Tyr-----ThrAsn 176  
 DB 1711 CAGGATGACATATTAGAAATTTCCAAAATGAAATCTGTGATTTAGCTGTTAGAAAAC 1770  
 QY 177 ValAspGluPheGly-----AlaSerPhePheAsnLysLeu 188  
 DB 1771 ATCAACAGATCCGACCATTAATCAACGTCACAAAGACCTGTTCAATCAACCAAGT 1830  
 QY 189 SerPheThrThrGly----- 193  
 DB 1831 CTATTGACGCTGGCAGGTCATTAACGCACTGCTAGATAAAGCGGCATATACCTTTC 1890  
 QY 194 -----LeuPheGlyTyrPglyIleLysArgAla 202  
 DB 1891 CGTGAATCGAAATGACCCGCTCTCAAGATTCCTCGGTGGTATAGCAAAACCGCA 1950  
 QY 203 LeuLysGlnIleIleArgSerAsnLeuProLeuAspIleGlyThrGlnHisSerValSer 222  
 DB 1951 CTA-----ATTGCTACTATATCGCTGCAAAAGGTAATCTTGAAAGAAACCTGCAGT 2001  
 QY 223 ArgLeuGlnHisIleThrSerSerTyrLysAspTyrMetAspThrGlnIleProAlaLeu 242  
 DB 2002 ACATTAGAGTAT---GCTTGACAGCCTAAACACTTAAGAAACAG-----CCGCAACTG 2052  
 QY 243 ProLysPheAlaLysArgPheSerLeuMet-----Val 253  
 DB 2053 GGTTCATTATTAAATGAAGATATTTGTTAAATAATTAATGAACTATGAAATGACAAAGATT 2112  
 QY 254 ValGlnArgLeuLeuAlaThrValAlaGly-----TyrValAspThrProTyr 270  
 DB 2113 AAATCGATTATCTCTCTACCAAGCTCAAGAGCAAGATATATATAGCCACATCACTACT 2172  
 QY 271 LysLysTyrTyrMetLysLeuLysAsnPheMetValAsnArgValPheIleProThrLys 290  
 DB 2173 AAAAATTTGAACAGCTGATTAGAAAGTTATAA---AATCAAGTT----- 2214  
 QY 291 LysPhePheAsnLysGluIleArgGluProSerLysAlaLeuLysGluLysValSerThr 310  
 DB 2215 -----CAAGATGATGAAGAAAGAAATTTGAAGTTTGACATCGAAGAAATGCAATTTG 2262  
 QY 311 AspThrLysAspLeuPheGluAsnLysIleGlyGlnIleThrValAspPhePheAsnLys 330  
 DB 2263 CTAGTAAAGATTAATTAATGAAGTCAAAA-----GAACTATTCATCTCAAAATGTC 2313  
 QY 331 GluIleArgAspProSerLysAlaLeu-----LysGlu 341  
 DB 2314 CAATATGAATCATTTGAAACTACATCATTTTAAGGCGACACACTAGATTAACACAT 2373  
 QY 342 LysValSerAsnAspAlaLysAspLeuPheGluAsnLysIleGlyGlnIleThr----- 359  
 DB 2374 AAAACGTAAATTTGAATATCCGAT---TTTAATTAACCAACTACAGAGTGTGAGAGTA 2430  
 QY 360 -----ValAspPhe----- 362  
 DB 2431 ATGCAATAGCCCTACATGATTACAAAAAAGAGAACCTTCAATCAAAAGTTTGA 2490  
 QY 363 -----IleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLysValSerThr 380  
 DB 2491 ATGCATATTACTAAAGAAATTAATAAATTTGAATCTACACTGTTTTCATTAATTAACACT 2550



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Oy 194 -----Leuphegilytrpgilylelysrghla 202
Db 1892 CGTGAATCGAAATGACCGCGCTTCAAGATTCCTGGTGTAAATACGAAACCGCA 1951
Oy 203 LeuylsglntlleleagserAsnLeuProleuspiilegylthrglunhisValSer 222
Db 1952 CTA-----ATTGCTACTATATCCCTGCAAGAGTAACCTTGTGAAGAAACCTGCAGT
Oy 223 ArgLeuGlnhisIlethrserserTyrLysAspTyrMetAspTyrGlnIleProAlaLeu 242
Db 2003 ACATTAAAGTAT---GCTTCAGAGCTAAACATTAAGAACAG-----CCCAACTG 2053
Oy 243 ProlysPheAlaLysArgPheSerLeuMet-----Val 253
Db 2054 GGTTCATTATTAATGAAGATATTGTGTTAAATATTAACATGAATTAAGCAAGAT 2113
Oy 254 ValGlnArgLeuAlaIlethValAlaIgly-----TyrValAspThrProTryptyr 270
Db 2114 AAATCCGATTTACTCTCTACAAAGTCCAAAGAGAAATATATAGCCCAAGATCACACTAC 2173
Oy 271 LysIleTryptyrMetLysLysAsnPhemeValAsnArgValPheIleProThrLys 290
Db 2174 AAAAATTTGAACACTGATTTGAAGAATTATAA---AATGAAGT----- 2215
Oy 291 LysPhePheAsnLysGluIleArgGluProSerLysAlaLeuLysGluLysValSerThr 310
Db 2216 -----CAAGAATGTAAGAAAGAAATGAAGATTGAACATCGAATCGAAATGATTC 2263
Oy 311 AsprhrLysAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPheAsnLys 330
Db 2264 CTACTAAAGATTAATGAAGTCAAAA-----GAACATATTCATTCACAAATTCG 2314
Oy 331 GluIleArgAspProSerLysAlaLeu-----LysGlu 341
Db 2315 CAATAGATTCATTTGAACACACATGATCATTTAAGGACACAACTGATTAACAGCAT 2374
Oy 342 LysValSerAsnAspAlaLysAspLeuPheGluAsnLysIleGlyGlnGlyThr----- 359
Db 2375 AAAAAGTAAATTTGAAGATTCGAT---TTTAATTAACAAACACAGAGTGTGCTGAGGTA 2431
Oy 360 -----ValAspPhe----- 362
Db 2432 ATGCAAAATGCCCTACATGATTCACAAAAAAGACAACTGTGACCTTAATCAAAAGTTTCAA 2491
Oy 363 -----IleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLysValSerThr 380
Db 2492 ATGCAATTTACTTAAGAAATTAATAAATTGAATCTACACTGTTTTCATTAACACT 2551
Oy 381 GlyAlaGluAsp-----LeuPheGluAsnLysIleGlyGlnGlyThrValAspPheIle 398
Db 2552 ATGCACAGAGAAAGTATTTCTTCAAGACCTAATATC---CAACCAAAATCTGATATGATC 2608
Oy 399 AsnAsnGluIleArgAspProSerLysAlaLeu----- 409
Db 2609 AAAAATGAGTACTGACTCTTATGACACCATGCAAGAAAAAGCTGAACATATGTACAAA 2668
Oy 410 -----IleArgLysValTyrThrGluAlaAspAspLeuPheGlu-----AsnLys 424
Db 2669 GACTGTGAGAAAGAAATTTAAAGCAATCTCTAAATCTTCAATGTTTCAATTACAGAAA 2728
Oy 425 IleGlyGlnGlyThrValAspPhe-----IleAsnLysGluIleArgAspProSerLys 442
Db 2729 ATTCACATTAATTAAGATGATTTCCAAAAATTTATTAATAATATATAGCGAGAAAT----- 2782
Oy 443 AlaLeuIleArgLysValSerThrGluAlaAspAsnLeu 455
Db 2783 -----CTTCTGATATTAGCGAAGAAATATACACATG 2815

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RESULT 17  
AAQ70102  
ID AAQ70102 standard: cdna to mRNA: 4766 BP.  
XX  
AC AAQ70102:

```

XX 22-SEP-1994 (first entry)
DT
XX Malarial PfEMP3 epitopic fragment clone p2b1.p12-1.
DE
XX
XX Plasmodium falciparum erythrocyte membrane protein, pfEMP3;
KW malaria; antigen; epitope; vaccine; anti-idiotypic antibody; ds.
XX
XX Plasmodium falciparum (Malayan Camp strain).
OS
XX
XX Key Location/Qualifiers
FH 3..4766
FT /*tag- a
FT /note- "partial coding region; does not include
FT initiation or termination codons"
XX
XX W09403604-A.
PN
XX
XX 17-FEB-1994.
PD
XX
XX 05-AUG-1993; 93WO-US07261.
PF
XX
XX 07-AUG-1992; 92US-0927531.
PR
XX
XX (SCHE ) SCHERING CORP.
PA
XX
XX Handumnetti SM, Howard RJ, Pasloske BL, Van Schravendijk MR.
PI
XX WPI: 1994-065693/08.
DR
XX P-PSDB: AAR46605.
DX
XX
XX New malaria antigen, pfEMP3 - used to isolate and produce prods.
PT for use in diagnosis, therapy and prevention of malarial
PT infection
XX
XX Claim 1; Page 72-79; 79pp; English.
PS
XX
XX The pfEMP3 malarial antigen is recognised by monoclonal antibody MAB
CC 12011. Nucleic acid sequences encoding part of the 315kd antigen,
CC have been isolated and sequenced. pfEMP3 is encoded on chromosome 2
CC of the P. falciparum genome and is thought to be associated with knob
CC formation and structure: malarial strains carrying deletions of the
CC gene coding for pfEMP3 exhibit a knobless phenotype.
CC
XX
SQ Sequence 4766 BP; 2404 A; 508 C; 863 G; 991 T; 0 other;
Alignment Scores:
Pred. No.: 0.212 Length: 4766
Score: 118.50 Matches: 57
Percent Similarity: 47.57% Conservative: 31
Best Local Similarity: 30.81% Mismatches: 66
Query Match: 5.02% Indels: 31
DB: Gaps: 14
US-09-807-459-2 (1-458) x AAQ70102 (1-4766)
Oy 294 AsnLysGluIleArg---GluProSerLysAlaLeuLysGluLysValSerThrAspThr 312
Db 3135 AATTAAGATTAACGAATTAAGATCTGAAGATTTAAAGAAATGCAACATAAAAT 3194
Oy 313 LysAspLeuPheGluAsnLysIleGlyGlnGly-----ThrValAspPheAsn 329
Db 3195 AAAAATTA---CAAAATTAAGATCTGAAGATTTAAAGAAATGCAAGATTTAAAAAT 3251
Oy 330 LysGluIleArgAspPro---SerLysAlaLeuLysGluLysValSerAsnAspAlaLys 348
Db 3252 AAAGATTAACAAATTAAGATCTGAAGATTTAAAGAAATGCAAGCAAAAAATTA 3311
Oy 349 AspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPheIleAsnAsnGluIleArg 368
Db 3312 GAATTA---CAAAATTAAGATCTGAAGATTTAAAGAA-----AATGCAAGATTTAAA 3362
Oy 369 AspProSerLysAlaLeuIleArgLysValSerThrGlyAlaGluAspLeuPheGlu--- 387

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Db 3363 -----AAATAAGATTTGCGAATAAGATCTGATGATTTAAAGAAATGCAAGCGTA 3416
Oy 368 -----AsnLysIleGlyGlyGly-----ThrValAspPheIle 398
Db 3417 AAAAATAAGATTTGCGAATAAGATCTGATGATTTAAAGAAATGCAAGATTTAA 3476
Oy 399 AsnAsnGluIleArgAspPro---SerLysAlaLeuIleArgLysValIleThrGluAla 417
Db 3477 AATTAAGATTTGCGAATAAGATCTGATGATTTAAAGAAATGCAAGATTTAA 3536
Oy 418 AspAspLeuPheGluAsnLysIleGlyGlyThrValAspPheIleAsnLysGluIle 437
Db 3537 GAT-----TTAAAGATTTGCGAATAAGATCTGATGATTTAAAGAAATGCAAGATG 3587
Oy 438 ArgAspProSerLysAlaLeuIleArgLysValIleThrGluAlaAsp----- 453
Db 3588 AAA-----AATTAAGATCTATTAACAAAGATTTCTTAATTAAGACATGAAGATTA 3641
Oy 454 AsnLeuLeuGluLys 458
Db 3642 GAACCTATTAAATTA 3656

RESULT 18
ID AAA70095
XX AAA70095 standard; DNA; 7326 BP.
AC
XX
XX AAA70095;
D7 07-NOV-2000 (first entry)
DE Plasmodium falciparum chromosome 2 related DNA sequence SEQ ID NO:228.
XX
XX Plasmodium falciparum; chromosome 2; human malaria parasite; vaccine;
KM antimalarial; malaria; protozoacide; infection; insecticide; ds.
XX
XX Plasmodium falciparum.
OS
XX WO200025728-A2.
PN
XX
XX 11-MAY-2000.
PD
XX
XX 05-NOV-1999; 99WO-US26796.
PE
XX
XX 05-NOV-1998; 98US-0107131.
PR
XX
PA (HOFF/) HOFFMAN S.
PA (GARU/) CARUCCI D.
PA (GARD/) GARDNER M.
PA (VENT/) VENTER J C.
XX
PI Hoffman S, Carucci D, Gardner M, Venter JC;
DR WPI: 2000-365347/31.
XX
XX Proteins encoded by chromosome 2 of the human malarial parasite,
PT Plasmodium falciparum, useful as antimalarial vaccines and in the
PT diagnosis of P.falciparum infection -
XX
XX
PS Disclosure: Page 454-456; 577pp; English.
XX
XX The present invention describes proteins and their fragments (I) encoded
CC by chromosome 2 of the human malarial parasite, Plasmodium falciparum.
CC Also described are: (I) nucleotide sequences (II) encoding (I); and (2)
CC vaccines against P. falciparum infection comprising (I) or (II).
CC (I) and (II) are useful for the development of vaccines against
CC P. falciparum infection. (I) and polyclonal antisera or a monoclonal
CC antibody raised to immunogens comprising the sequences of (I), are
CC useful in the detection of infection with P. falciparum. Furthermore,
CC (I) (especially when they are rifins or secreted or membrane proteins)
CC can aid the identification of drugs to treat or prevent P. falciparum
CC infection, or they can be used to identify drug resistance in
CC P. falciparum. Sequencing of the Plasmodium chromosome 2 and the

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CC subsequent identification of proteins encoded by it will help to expand
CC our understanding of parasite biology, a process hampered by the
CC complexity of the parasitic lifecycle, and provide new targets for
CC vaccine and drug development. Parasite resistance to drugs and mosquito
CC resistance to insecticides have led to a resurgence of malaria in many
CC parts of the world, and there is a pressing need for vaccines and new
CC drugs. AAA70078 to AAA70287 and AAB18144 to AAB18352 represent nucleotide
CC and protein sequences given in the present invention, but which are not
CC specifically mentioned within the specification.
XX
SQ Sequence 7326 BP; 3588 A; 868 C; 1321 G; 1549 T; 0 other;
XX
Alignment Scores:
Pred. No.: 0.377 Length: 7326
Score: 118.50 Matches: 58
Percent Similarity: 48.658 Conservative: 32
Best Local Similarity: 31.35% Mismatches: 64
Query Match: 5.02% Indels: 31
DB: Gaps: 14

US-09-807-459-2 (1-458) x AAA70095 (1-7326)
Oy 294 AsnLysGluIleArg---GluProSerLysAlaLeuLysGluLysValSerThrAspThr 312
Db 3361 AATTAAGATTTGCGAATAAGATCTGGAAGATTTAAAGAAATGCAAGATTTAA 3420
Oy 313 LysAspLeuPheGluAsnLysIleGlyGlyGly-----ThrValAspPheIleAsn 329
Db 3421 AAAAGATTTA---CAAAATTAAGATCTGGAAGATTTAAAGAAATGCAAGATTTAA 3477
Oy 330 LysGluIleArgAspPro---SerLysAlaLeuLysGluLysValSerAsnAspAlaLys 348
Db 3478 AAAAGATTTGCGAATAAGATCTGGAAGATTTAAAGAAATGCAAGATTTAA 3537
Oy 349 AspLeuPheGluAsnLysIleGlyGlyGlyThrValAspPheIleAsnAsnGluIleArg 368
Db 3538 GAATTA---CAAAATTAAGATCTGGAAGATTTAAAGAAATGCAAGATTTAA 3588
Oy 369 AspProSerLysAlaLeuIleArgLysValSerThrGluAlaGluAspLeuPheGlu--- 387
Db 3589 -----AATTAAGATTTGCGAATAAGATCTGGAAGATTTAAAGAAATGCAAGATTA 3642
Oy 388 -----AsnLysIleGlyGlyGly-----ThrValAspPheIle 398
Db 3643 AAAAATAAGATTTGCGAATAAGATCTGGAAGATTTAAAGAAATGCAAGATTTAA 3702
Oy 399 AsnAsnGluIleArgAspPro---SerLysAlaLeuIleArgLysValIleThrGluAla 417
Db 3703 AATTAAGATTTGCGAATAAGATCTGGAAGATTTAAAGAAATGCAAGATTTAA 3759
Oy 418 AspAspLeuPheGluAsnLysIleGlyGlyGlyThrValAspPheIleAsnLysGluIle 437
Db 3760 AATGATTTAAAGATTAAGATTAATTT-----CAAAATTAAGATTTATCTAATAAGACATG 3813
Oy 438 ArgAspProSerLysAlaLeuIleArgLysValSerThrGluAlaAsp----- 453
Db 3814 AAA-----AATTAAGATCTATTAACAAAGATTTCTTAATTAAGACATGAAGATTA 3867
Oy 454 AsnLeuLeuGluLys 458
Db 3868 GAACCTATTAAATTA 3882

RESULT 19
AAAB1514/C
ID AAAB1514 standard; DNA; 33303 BP.
XX
XX
XX AAAB1514;
AC
XX
DT 04-DEC-2000 (first entry)
XX
XX N. meningitidis partial DNA sequence gnm_61 SEQ ID NO:61.
DE
XX
XX Neisseria meningitidis; Neisseria gonorrhoeae; genome; immunogenic;

```

antigen; vaccine; diagnosis; infection; antibacterial; identification;  
KM Meningococcus B; MenB; ds.  
OS Neisseria meningitidis.  
PN MO200022430-A2.  
XX 20-APR-2000.  
XX 08-OCT-1999; 99MO-US23573.  
XX 09-OCT-1998; 98US-0103794.  
PR 30-APR-1999; 99US-0132068.  
XX (CHIR ) CHIRON CORP.  
PA Frazer CM, Hickey E, Peterson J, Tettelein H, Venter JC,  
PI Maignan V, Galeotti C, Mora M, Ratcli G, Scarselli M, Scarlato V;  
PI Rappuoli R, Pizza M;  
XX WPI; 2000-318079/27.  
DR  
XX Isolated nucleotide sequences of Neisseria meningitidis which can be  
PT used in the diagnosis and treatment of N. meningitidis infection and  
PT other Neisserial infections, for example, N.gonorrhoea -  
XX  
XX Claim 7; Page 1375-1385; 1760pp; English.  
CC The present invention describes methods of obtaining immunogenic  
CC proteins from Neisseria genomic sequences. AAA81453 to AAA82414  
CC represent specifically claimed Neisseria meningitidis genomic DNA  
CC sequences: AAA81260 to AAA81303 and AAB25620 to AAB25663 represent  
CC Neisseria DNA sequences and their corresponding proteins: AAA81254 to  
CC AAA81259 and AAA81304 to AAA81321 represent PCR primers used in the  
CC isolation of Neisseria meningitidis DNA sequences; and AAA81322 to  
CC AAA81352 represent Neisseria meningitidis MenB polynucleotide ORF  
CC sequences, which are all used in the exemplification of the present  
CC invention. The nucleic acid sequences, protein sequences, and antibodies  
CC against them, can be used in the manufacture of a composition. The  
CC composition can be used as a medicament (or in the manufacture of a  
CC medicament) for treating, preventing or diagnosing infection due to  
CC Neisserial bacteria. For example, some of the identified proteins could  
CC be components of vaccines against Meningococcus B; against all serotypes;  
CC and/or against all pathogenic Neisseriae. Identification of sequences  
CC from the bacterium will also facilitate production of biological probes,  
CC particularly organism-specific probes. Attempts to make efficacious  
CC Meningococcus B vaccines have failed mainly due to antigen tolerance.  
CC Multivalent vaccines have also been tried but none have successfully  
CC overcome antigenic variability. The provision of further, complete  
CC sequences may provide an opportunity to identify secreted or surface  
CC exposed proteins that may be presumed targets for the immune system and  
CC which are not antigenically variable or at least more conserved than  
CC other more variable regions.  
XX  
SQ Sequence 33303 BP; 7919 A; 8691 C; 8422 G; 8266 T; 5 other:  
  
Alignment Scores:  
Pred. No.: 4 Length: 33303  
Score: 117.00 Matches: 81  
Percent Similarity: 35.66% Conservative: 62  
Best Local Similarity: 20.20% Mismatches: 150  
Query Match: 4.96% Indels: 108  
DB: 21 Gaps: 19  
  
US-09-807-459-2 (1-458) x AAA81514 (1-33303)  
  
OY 94 LeuProAspProTyrGlnLeuGluAlaAlaPheIleLeuPheLysGluSerAspAlaasn 113  
DB 33123 TTGGGTGACCGCTTATGCGAGATGCTGCAATGTCGTAAGAGCAAACTTAAT 33064  
OY 114 ProLAsnSerThrGluLysArgPheMetArgPheMetArgGlyLysasnHisser 133  
DB 33063 GGTTCGAAT-----TTATGATG-----AAAAAGGT----- 33037

OY 134 TyrPheHisAspLeuValPheAsnLeuGluLysAsnValThrArgAspAlaAspAla 153  
DB 33036 -----GTGCAAAACCTATAGCATGATACGGTCGGTAA-----AAG 33001  
OY 154 ThrAspIleGluAsnPheAlaSerArgTyrLeuTyrMetAlaThrLeuTyrTyrLysThr 173  
DB 33000 ACCCGTTAGAGAAATTGATCGG-----GTTGCACGTGCAACATTCAGCGCA 32953  
OY 174 TyrThr -----AsnValAspGluPheGlyAla 182  
DB 32952 TATGGCGCTTAATTATCAAAATAGTAGATTACCAATACAGTAATGGAATTGACAGA 32893  
OY 183 SerPheAsnLysLeuSerPheThrThrGluLeuPheGlyTyrGlyLysArgAla 202  
DB 32892 AGTACTATTAAGCCGTTACCGATAAT-----GGCGTTTCTCCAGTCAGCAGCT 32845  
OY 203 LeuLysGluIleIleArgSerAsnLeuPro-----LeuAsp 214  
DB 32844 ATGTATTAGTTAATTATCGTTCACTCCGATATGCGGATGTTATTTGGCATTAAGCT 32785  
OY 215 IleGlyThrGlu-----HisSerValSerArgLeuGlnHisIleThrSerSer 230  
DB 32784 TTGGGATAGAACCGCAACGATATCCACATAGCAACAGCATTAATATCCGACGATAC 32725  
OY 231 TyrLysAspTyrMetAspThrGlnIleProAlaLeuProLys----- 244  
DB 32724 GAAGGCGATATAGAACACATTAATATCTGCTTTAGATTAAGATTTGATGATCTTTT 32665  
OY 245 PheAlaLysArgPheSerLeuMetValAlaGlnArgLeuAlaThrValAlaGly--- 263  
DB 32664 AAAGAGAACGCAATTTACTTTTTCATCATCTGTGATGATGATGTAACAAAGTTAGCTT 32605  
OY 264 ---TyrValAspThrProTyrTyrLys-----LysTrp-----TyrMetLys 276  
DB 32604 GATATTACAATATAGATGCTTGGCAAAAATTTGGAGGTTGGGTAATGGATATACATAT 32545  
OY 277 LeuLysAsnPheMetValAsnArgValPheIleProThrLysLysPheAsnLysGlu 296  
DB 32544 TTATATAAAGGTTGTTAAAGAGAGAGCTGACATGATTAATGATGTTAATTAATAC 32485  
OY 297 IleArgGluProSerLysAlaLeuLysGluLysValSer----- 309  
DB 32484 ATCAAGCAAGAGAAATGAAGCTTTTAAATGAATCAATACCTGCTTCATGATATGAAA 32425  
OY 310 -----ThrAspThr 312  
DB 32424 GCTGCTGCAAGCAATTTGAGATGACTTAATATACAGCTGGAATATATCTCAGCTGAGCT 32365  
OY 313 LysAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPhePheAsnLysGluIle 332  
DB 32364 GCCCAATATACTTAATAGATATAGTACATACACTCAAGCAATAGAAAAGGTTCTC 32305  
OY 333 ArgAspProSerLysAlaLeuLysGluLysValSerAsnAspAlaLysAspPheGlu 352  
DB 32304 AAA---GCCATTAAAGAAATTTGCTGAAAAAATGCTGCTCCGATTTGGCTAC 32248  
OY 353 AsnLysIleGlyGlnGlyThrValAspPheIleAsnAsnGlnIleArgAspProSerLys 372  
DB 32247 GGTTC-----GCAGAGAAACCTAAA 32227  
OY 373 AlaLeuIleArgLysValSerThrGlyAlaGluAspLeuPheGluAsnLysIleGlyLys 392  
DB 32226 CAAGTAGTGAAGATTTGCTCAACCCCAAGCAAGCATCGAAAT-----GCCAAA 32173  
OY 393 GlyThrValAspPheIleAsnAsnGlnIleArgAspProSerLysAlaLeu-----Ile 410  
DB 32172 TCCACAGCCGAGAAAGCGCTCCACACCTCGAGAAATTTTAAAGCGTTTGGCCAGTTT 32113  
OY 411 ArgLysValTyrThrGluAlaAspAspLeuPheGluAsnLysIleGlyGlnGlyThrVal 430  
DB 32112 AAAGATCTGGCGGAAAAATTTAGAGATCTGTTCCCAATTCG-----GAAGGCTGATC 32059

Oy 431 Asp 431  
 |||  
 Db 32058 GNT 32056  
 RESULT 20  
 ID AAF21610 standard; DNA: 349980 BP.  
 XX AAF21610  
 AC AAF21610;  
 DT 13-MAR-2001 (first entry)  
 XX  
 DE Neisseria meningitidis B nucleotide sequence SEQ ID NO:111.  
 XX  
 KW Neisseria meningitidis; Neisseria gonorrhoeae; immunogenic; vaccine;  
 KW diagnosis; antigen; detection; infection; gene therapy; antibacterial;  
 ds.  
 XX  
 OS Neisseria meningitidis.  
 XX  
 PN WO200006791-A1.  
 XX  
 PD 09-NOV-2000.  
 XX  
 PF 08-MAR-2000; 2000WO-US05928.  
 XX  
 PR 30-APR-1999; 99US-0132068.  
 PR 08-OCT-1999; 99WO-US23573.  
 PR 28-FEB-2000; 2000GB-0004695.  
 XX  
 PA (CHIR ) CHIRON CORP.  
 PA (GENO-) INST GENOMIC RES.  
 XX  
 PI Piazza M, Hickey E, Peterson J, Tettelin H, Venter JC, Maignani V;  
 PI Galeotti C, Mora M, Ratti G, Scarselli M, Scariato V, Rappunli R;  
 PI Frazer CM, Grandi G;  
 XX  
 DR WPI: 2000-647603/62.  
 XX  
 XX Neisseria meningitidis B full length genome sequence and open reading  
 PT frames are used to detect, treat and prevent Neisserial infections -  
 PS  
 PS Claim 7; Appendix A: 692pp: English.  
 XX  
 CC The present invention describes the full length genome of  
 CC Neisseria meningitidis B (NMB). The sequences in AAF21544 and AAF21607  
 CC to AAF21613 represent fragments of the NMB genomic sequence, as the  
 CC sequence was too long to go in a record on its own it was split into 8  
 CC sequences which overlap each other at the beginning and end of each  
 CC sequence by 49980 bp (i.e. the last 49980 bp of AAF21544 is repeated at  
 CC the beginning of AAF21607, the last 49980 bp of AAF21607 are repeated at  
 CC the beginning of AAF21608, and so on). AAF21545 to AAF21588 encode the  
 CC Neisseria proteins given in AAB58550 to AAB58593, and AAF21589 to  
 CC AAF21606 represent PCR primers which are used in the exemplification of  
 CC the present invention. The NMB genome and fragments from it have  
 CC antibacterial activity, and can be used in vaccines and gene therapy.  
 CC Neisseria nucleic acids, proteins and/or antibodies which binds to the  
 CC proteins can be used in compositions for treating or preventing infection  
 CC due to Neisserial bacteria or as a diagnostic reagent for detecting the  
 CC presence of Neisserial bacteria or of antibodies raised to Neisserial  
 CC bacteria. Computers, computer memory, computer storage medium or computer  
 CC databases can be used in a search to identify open reading frames (ORFs)  
 CC or coding sequences within the NMB genome. The DNA sequences provide  
 CC further opportunities to find antigenic or immunogenic proteins which are  
 CC more effective in vaccines than the outer membrane proteins currently  
 CC used.  
 XX  
 XX Sequence 349980 BP; 86771 A; 92803 C; 86340 G; 84066 T; 0 other;  
 Alignment Scores:  
 Pred. NO.: 92.3 Length: 349980  
 Score: 117.00 Matches: 81  
 Percent Similarity: 35.668 Conservative: 62

Best Local Similarity: 20.20%		Mismatches: 150	
Query Match: 4.96%		Indels: 108	
DB:	21	Gaps: 19	
US-09-807-459-2 (1-458) x AAF21610 (1-349980)			
OY	94	LeuProAspProGlyGlnLeuGluAlaIalaPheIleuPheLysGlnSerAspAlaAsn	113
Db	243690	TTGGGTGACCGCTTATGGCGATGATGCTGCTCAATTCTCGTAAGATGCAACTTAAT	243749
OY	114	ProAlaAsnSerThrGlnLysArgPheThrMetArgPheArgArgLysAsnHisSer	133
Db	243750	GGTTTGGAT-----TTATGATG-----AAAAAGGT-----	243776
OY	134	TyrPheHisAspLeuValPheAsnLeuGluLysAsnValThrArgAspAlaAspAla	153
Db	243777	-----GTGGAAACACCTATGGATGATACGGCGGTAA-----AAG	243812
OY	154	ThrAspIleGluAsnPheAlaSerArgTyrLeuTyrMetIaThrLeuTyrThr	173
Db	243813	ACCCGTTTGAAGAAATTTGATCG-----GTGCACATGCACATTTTCAGGCAA	243860
OY	174	TyrThr-----AsnValAspGluPheGlyAla	182
Db	243861	TATGCGCCTCAATTAATCAAAATAATGGTGATATTACCAATACTACTAGTAATTGAGACA	243920
OY	183	SerPhePheAsnLysLeuSerPheThrThrGlyLeuPheGlyTrpGlyIleLysArgAla	202
Db	243921	AGTTACTATTAAGCCGTTACCGATAAT-----GCCGTTTCTTCAGTCAGCT	243968
OY	203	LeuLysGlnIleIleArgSerAsnLeuPro-----LeuAsp	214
Db	243969	ATTGATTTAGTTATTAATCTGTCCTCCGATATGGCGAGTGTATTGGCATTAAGT	244028
OY	215	IleGlyThrGlu-----HisSerValSerArgLeuGlnHisIleThrSerSer	230
Db	244029	TTGGGATAGAGAACCCGAACGATCTCCACATAGACAGACGTAATTAATCCGAACGGTAGC	244088
OY	231	TyrLysAspTyrMetAspThrGlnIleProAlaLeuProLys-----	244
Db	244089	GAAGAGGTATATAGAAACGACTATATCTGCTTTAGATGAAGATTGATGATCTTTT	244148
OY	245	PheAlaLysArgPheSerLeuMetValValGlnArgLeuLeuAlaThrValAlaGly--	263
Db	244149	AAAGAGACGATTTTACTTTTTCACATCTGTGATGATGATGATGACAAAGTTAGTGTT	244208
OY	264	--TyrValAspThrProTrpTyrLys-----LysTrp-----TyrMetLys	276
Db	244209	GAATATACATATAGATGCTTGGCCAAAAAATTTGGAGTTGGGTAAATGCGATATCATGAT	244268
OY	277	LeuLysAsnPheMetValAsnArgValPheIleProThrLysLysPhePheAsnLysGlu	286
Db	244269	TTATATATAAAGTGTGTATAAAAGAGAGCGACGTGGAATATTGAGATCGTTAATAATAAC	244328
OY	297	IleArgGluProSerLysAlaLeuLysGluLysValSer-----	309
Db	244329	ATCAAGCAAGAAATGAAGCTTTTAAAAATGAAATCATATACCTGTTCATGATATGAAA	244388
OY	310	-----ThrAspThr	312
Db	244389	GCTGCTGGCAAGAAATTTGGAGATGACTTAATAATACAGATGCGAATAATCTCACACAGCT	244448
OY	313	LysAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPhePheAsnLysGluIle	332
Db	244449	GCCGAATATATCTATATATGACATAGTACACATATCTAGTACAGAAATGAAAAAGGTGTC	244508
OY	333	ArgAspProSerLysAlaLeuLysGluLysValSerAsnAspAlaLysAspLeuPheGlu	352
Db	244509	AAA--GCCATTAATAAGATTGTCTGAAAAAATGAAAAATGCTGCTCCGATTGGCTGAC	244565
OY	353	AsnLysIleGlyGlnGlyThrValAspPheIleAsnAsnGlnIleArgAspProSerLys	372
Db	244566	GATTCA-----GCAGAGAAAGCTAAA	244586







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Db 3996 AAMACAGCATTCAGAGAGCGGCTTGCTTACGACGCTCCAGCATTCATGAGTGT 4055
Oy 221 lserArgLeuGlnHisIleThrSerSerTyrLysAspTyrMetAspThrGlnIleProAl 241
    |||||
Db 4056 GAGCAGCAATTCAGAGCGCTGCTCCAGCTTACAGTACACAGAGAT----- 4107
Oy 241 aleuProLysPheAlaLysArgPheSerLeuMetValValGlnArgLeuAlaIleThrVa 261
    |||||
Db 4108 -----CAGCAGCAGCTTCAGCAGATT 4127
Oy 261 lAlaGlyTyrValAspThrProTyrTyrLysTyrIleTyrMetLysLeuLysAsnPheMe 281
    :
Db 4128 GGGCTCCACCGACGACGCT----- 4149
Oy 281 tValAsnArgValPheIleProThrLysPhePheAsnLysGlnIleArgLupProse 301
    |||||
Db 4150 -----AAGGAA---ACCAGCCCTTC 4166
Oy 301 rLysAlaLeuLysGlnLysValSerThrAspThrLysAspLeuPheGlnAsnLysIleG1 321
    |||||
Db 4167 GGGTCCCTTAAGCGAACCTACTCCAGTCCAGCAAAATCAGCTGCCGGAATAATTCa 4226
Oy 321 yGlnGlyThrValAspPheAsnLysGlnIleArg-----AspProSerLysAl 338
    |||||
Db 4227 ACAG-----TTATGAGACAAATCAACGCTCAACTACAGAAAGCGAGC 4271
Oy 338 aLeuLysGlnLysValSerAsnAspAlaLys---AspLeuPheGlnAsnLysIleG1 357
    |||||
Db 4272 CGAAAGAGAGAGAGCTGCGACCAACGAGCGGAATACCTTCAAATCTGCAGTTTCA 4331
Oy 357 nGlyThrValAspPheIle-----AsnAsnGlnIleArgAspProSer 371
    |||||
Db 4332 AGGCGCAGCAGGAATGTGTCTACACAGAAATTAACATTCTCGCGCAAGC 4383
RESULT 23
AAV84691 standard; DNA; 10478 BP.
XX
AC AAV84691:
XX
DT 12-APR-1999 (first entry)
XX
DE Arabidopsis ESD4 (early short days 4) gene.
XX
KW ESD4 gene: early short days 4; flowering; transgenic plant; ss.
XX
OS Arabidopsis thaliana.
XX
FH key Location/Qualifiers
FT CDS 3490..6495
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    /*note= "contains introns"
    3490..4239
FT exon /*tag= b
    /*number= 1
    4240..4535
FT intron /*tag= c
    /*number= 1
    4536..4612
FT exon /*tag= d
    /*number= 2
    4613..4689
FT intron /*tag= e
    /*number= 2
    4690..4780
FT exon /*tag= f
    /*number= 3
    4781..4851
FT intron /*tag= g
    /*number= 3
    4852..4950
FT exon /*tag= h
    /*number= 4
FT

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```

FT intron 4951..5247
FT /*tag= i
FT /*number= 4
FT exon 5248..5337
FT /*tag= j
FT /*number= 5
FT intron 5338..5435
FT /*tag= k
FT exon 5436..5555
FT /*tag= l
FT intron 5556..5802
FT /*tag= m
FT exon 5803..5910
FT /*tag= n
FT intron 5911..6198
FT /*tag= o
FT exon 6199..6271
FT /*tag= p
FT intron 6272..6432
FT /*tag= q
FT exon 6433..6495
FT /*tag= r
FT /*number= 9
FT /*tag= s
FT /*note= "this region is from ecotype Landsberg
    erecta, the remainder of the sequence is
    from ecotype Columbia"
FT mutation 2757..3518
    /*tag= t
    /*note= "this region is deleted in the eds4 mutant"
FT
XX
PN W09856918-A1.
XX
PD 17-DEC-1998.
XX
PF 12-JUN-1998; 98WO-GB01714.
XX
PR 13-JUN-1997; 97GB-0012415.
XX
PA (PLAN-) PLANT BIOSCIENCE LTD.
XX
PI Coupland GW;
PI
DR WPI: 1999-070324/06.
DR P-PSDB: AAM86184.
XX
PT New nucleic acid of the Arabidopsis thaliana ESD4 gene - used for
PT regulating flowering time in plants
XX
PS Example 1; Fig 2; 73pp; English.
XX
CC This is the nucleotide sequence of the early short days 4 (ESD4)
CC gene of Arabidopsis thaliana ecotype Columbia (with part of the
CC sequence from an ecotype Landsberg erecta clone) that plays a role
CC in the regulation of flowering time. Plants carrying a recessive
CC mutation affecting the ESD4 gene flower earlier than their
CC wild-types under long and short days. The eds4 mutant was
CC identified in a population of gamma-ray mutagenised plants. The
CC ESD4 gene was then identified by complementation and sequencing of
CC the mutant allele. Regulation of ESD4 gene expression is used to
CC alter (advance or delay) the time of flowering of transgenic
CC plants. Overexpression delays flowering while underexpression
CC promotes it, e.g. to ensure simultaneous flowering of both parents
CC in hybrid production, or to regulate flowering according to
CC climatic conditions. ESD4 nucleic acids can also be used as

```

CC primers or probes to isolate homologous sequences in other plants  
 CC (see also AAV84692-93).

SO Sequence 10478 BP; 3195 A; 1770 C; 1986 G; 3527 T; 0 other;

# Alignment Scores:

Pred. No.:	1.89	Length:	10478
Score:	113.50	Matches:	99
Percent Similarity:	38.29%	Conservative:	89
Best Local Similarity:	20.16%	Mismatches:	190
Query Match:	4.81%	Indels:	113
DB:	20	Gaps:	27

US-09-807-459-2 (1-458) x AAV84691 (1-10478)

```

OY 14 LeuLeuAlaAlaSerGluSerValAspSerAlaAlaAsnAlaTyrMetIleAsnSerAsp 33
    :::::|||||::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 8178 ATCTTGCAGGAATTCGAGAGAGAGAGAGATGATGCTGACTATCATCACTCGTCACGGAT 8237
OY 34 MetSerAspTyrLeuSerAlaValSerAspAsnPheAlaGluArgIle----- 49
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 8238 CTCAATCTCTACTTGATACAGACGATGATGAAATTCATCATCGATTCTACGGTAGTCTC 8297
OY 50 -----Cys-----SergIleValPro 54
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 8298 TCCTCAATGCGCTTCTCGTTTCTCCGCTGTAATCCGCCGCCGATGATTCGCCGTTGAA 8357
OY 55 LysGlySerAsnCysSerAlaSerValSerAlaTyrMetSerArgCysAlaLysGlnAsp 74
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 8358 TCAGCCCTCTTGCTATTTTATGACCCGACGCTATTCACAGCTCTCTCTTCCACGAAAT 8417
OY 75 CysLeuThrIleGlnSerLeuIleTyrProLeuGlnAlaLysTyrGlnProLeuThrIleu 94
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 8418 TGC-----CCCGTTTCACTTCTCTCTCTCCGATTGCGTTT 8453
OY 95 ProAspProTyrGlnLeuGlnAlaAlaPheIleLeuPheLysGlnSerAspAlaAsnPro 114
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 8454 CTTCCTACTTGGCTCATTCGCGCTACCTCAAAACGCCGATGATTCGCCGCTCAA 8513
OY 115 AlaAsnSerThrGlnLysArgPheThrMetArgPheArgArgGlyLysAsnHisSerTyr 134
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 8514 GCGAGTAATTTGCGACGAGAGAAAGCAATGAAGAAGAGAGACGCTGT----- 8564
OY 135 PheHisAspLeuValPheAsnLeuGlnLysAsnVal---ThrArgAspAlaAspAla 153
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 8565 -----AAGAGGATTTAGCGTATGACGATGACGATGCGGAAGG 8597
OY 154 ThrAspIleGluAsnPheAlaSerArgTyrLeuTyrMetAlaThrLeuTyrTyrLysThr 173
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 8598 ACTGAAGAAGGTGATTTGATCGCAATGATGATTATATAGTG----- 8639
OY 174 TyrThrAsnValAspGluPheGlyAla-----SerPhePheAsnLysLeuSerPheThr 191
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 8640 -----TTAGAGAACCTTGGCTCGGCTTCTGAGTTTGTTCATGATAGATTAGTTTCTT 8690
OY 192 ThrGlyLeuPheGlyTyrGlyIleLysArgAlaLeuLysGlnIleIleArgSerAsnLeu 211
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 8691 GATAGTTTG-----AAATCTTTGGTACAAACTGTG---AGTGAGAT 8729
OY 212 ProLeuAspIleGlyThrGluHisSer-----ValSerArgLeuGlnHisIle 227
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 8730 CCTTTA---TTGGCGTGTGACACACTGTGAGTTTGAATTATGATGATGATGGAATG 8786
OY 228 ThrSerSerTyr-----LysAspTyrMetAspThrGlnIleProAla 241
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 8787 TGTGGGAAGATTCTGGGAGAGAGTGTGAATTCGTGACATGAGATAGGCACTCACTGCT 8846
OY 242 LeuPro-----LysPheAlaLysArg 248
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 8847 GCTGAGATTTCAGAGCTTTTGACACCGTTCTTATGCGTTTACATCAACGCAAGAAAT 8906
OY 249 PheSerLeuMetValValGlnArgLeuLeuAlaThrValAlaGlyTyrValAspThrPro 268
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|

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DB 8907 TTTGGCTGGCATTTGTGTCAAGAAATGATGAGTTTGCT-----AAAGATAACCT 8960
OY 269 TrpTyrLysLysTrpTyrMetLysLysAsnPheMetValAsnArgValhellePro 288
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 8961 GAATTGAAAAAAGTTGTCTTAATTTGCCCTAAGTTTCTGCTCATTAAGCA-----CCT 9014
OY 289 ThrLysLys-----PhePheAsnLysGlnIleArgGluProSerLysAlaLeu 304
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 9015 GAGAAAGCCGACCCCGCTGATTTGACAGAGAGCGGCTGTGAGATTATAAGCAAG 9074
OY 305 LysGlnLysValSerThrAspThrLysAspLeuPheGlnLysLysIleGlyGlnGlyThr 324
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 9075 GAGTTGAGGGCCCAATCAGAGTTTGTTGAT---TTTGAATGAGATGCTCAAGCAAG 9131
OY 325 ValAspPhePheAsnLysGlnLysLeuArgAspProSerLysAlaLeuLysLysValSer 344
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 9132 TCTAATTTTAGAGTATTAGCTGTT---GATATTATACCTCTGTGTGTAAGCTCATTAAGA 9188
OY 345 AsnAspAlaLysAspLeu-----PheGlnAsnLysIleGlyGlnGlyThr 359
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 9189 AACCTCTAGAGATATTATGTTTCAGAGATGAGGTTGAAGATTCGTGGGCTTGCTGT 9248
OY 360 ValAspPheIleAsnAsnGlnLysLeuArgAspProSerLysAlaLeuLysValSer 379
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 9249 ATTGATGCTTTAGTTTCAGCGGCTTTCAGACAGCAGC---GCTTTGATTTAGA----- 9296
OY 380 ThrGlyAlaGlnAspLeuPheGlnLysLysIleGlyGlnGlyThrValAspPheIleAsn 399
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 9297 -----GCTCGAGCTTTG---TCCAACTTGCTCAAA---GTTGTGAGTCTTGTCT 9341
OY 400 AsnGlnIleArgAspProSerLysAlaLeuIleArgLysValTyrThrGlnAlaAspAsp 419
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 9342 GGTGATGAAGG-----AGTAGTCGATCCGAAACAGCCCTTGCTTAACGCTGAG 9395
OY 420 LeuPheGlnAsnLysIleGlyGlnGlyThrVal---AspPheIleAsnLysGlnLysLeuArg 438
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 9396 ACTTCAGAG-----GGAAGAGTGCAGTAACGACCTTTTGAAGAAAGATGTGTG 9446
OY 439 AspProSerLysAlaLeuLysArgLysValSer 449
    ::::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
DB 9447 GAT---GAGAGCGCGCTGTAAAGAGACGACT 9476

```

## RESULT 24

AA576165 ID AAS76165 standard; cDNA; 4282 BP.

AC AAS76165;

DT 13-FEB-2002 (first entry)

DE DNA encoding novel human diagnostic protein #11969.

KW Human; chromosome mapping; gene mapping; gene therapy; forensic;

KW food supplement; medical imaging; diagnostic; genetic disorder; ss.

OS Homo sapiens.

PN WO200175067-A2.

PD 11-OCT-2001.

PF 30-MAR-2001; 2001WO-US08631.

PR 31-MAR-2000; 2000US-0540217.

PR 23-AUG-2000; 2000US-0649167.

PA (HYSE-) HYSEQ INC.

PI Drimanac RT, Liu C, Tang YT;

DR WPT: 2001-639362/73.

DR P-PSDB; ABG11978.

XX

PT New isolated polynucleotide and encoded polypeptides, useful in diagnostics, forensics, gene mapping, identification of mutations responsible for genetic disorders or other traits and to assess biodiversity -

PS Claim 1; SEQ ID No 11969; 103pp; English.  
xx

The invention relates to isolated polynucleotide (I) and polypeptide (II) sequences. (I) is useful as hybridisation probes, polymerase chain reaction (PCR) primers, oligomers, and for chromosome and gene mapping, and in recombinant production of (II). The polynucleotides are also used in diagnostics as expressed sequence tags for identifying expressed genes. (II) is useful in gene therapy technique to restore normal activity of (II) or to treat disease states involving (II). (II) is useful for generating antibodies against it, detecting or quantitating a polypeptide in tissue, as molecular weight markers and as a food supplement. (II) and its binding partners are useful in medical imaging of sites expressing (II). (I) and (II) are useful for treating disorders involving aberrant protein expression or biological activity. The polypeptide and polynucleotide sequences have applications in diagnostics, forensics, gene mapping, identification of mutations responsible for genetic disorders or other traits to assess biodiversity and to produce other types of data and products dependent on DNA and amino acid sequences. A5654197-A5934564 represent novel human diagnostic coding sequences of the invention. Note: The sequence data for this patent did not appear in the printed specification, but was obtained in electronic format directly from WIPO at [http://wipo.int/pub/published\\_pcc\\_sequences](http://wipo.int/pub/published_pcc_sequences).

50 Sequence 4282 BP; 1735 A; 915 C; 751 G; 881 T; 0 other;

**Alignment Scores:**

Pred. No.:	0.64	Length:	4282
Score:	113.00	Matches:	109
Percent Similarity:	33.04%	Conservative:	52
Best Local Similarity:	23.04%	Mismatches:	166
Query Match:	4.79%	Indels:	146
DB:	23	Gaps:	24

US-09-807-459-2 (1-458) x AAS76165 (1-4282)

OY	55	lyscGlySerMetGlySerMetIleSerValSerAlaTyrMetSerArgCysAlaLysGlnSer	74
		::: :	
Db	1725	AAAGGGATTTCATTTCAGAAAGAGCAAGTCAAAATTTGCTCCGTGTTGCGAGCAACATGCT	1784
OY	75	CysLeuThrLeuGln-----SerLeuLysTyr-ProLeuGlnAlaLys-----	88
		::: :	
Db	1785	TGTTTATCTAGAAAAACCCCATGCTGCACGCCCAAAATCTCTTAAGCTGATGAACACATT	1844
OY	89	TyrGlnProLeuThrLeuProLysPro-TyrGlnLeuGlnAlaIleAlaPheIleLeuPheLys	108
		::: :	
Db	1845	CACCAAAAGTCTCAGGATACAAAATCAATGATACAAAATACAAAGCATCTTTATAC----	1895
OY	108	sgLysSerPalaAsnProAlaLysSerThrGlnLysArgPheThrMetArgPheArgArg	128
		::: :	
Db	1900	-----ACCAATACACACAGCAAA-----	1917
OY	128	gGlyLysAsnHisSerTyrPheHisAspLeuValPheAsnLeuLeuGlyLysAsnVal--	147
		::: :                    :::	
Db	1918	-----CAAGAGGCCAATCATGATCAATCCCATTCACAACTGCTTCACAAAGCAATAAA	1972
OY	148	-----ThrArgSerPalaAsnAlaSerThrAspIleGluAsnPheAlaSe	161
Db	1974	ATACCTAGCAATCCAACTTACAAAGGATGTGAAGGACSTCTTCAAAGAGAACTACAAACC	2033
OY	161	r-----	161
Db	2034	ACTGTCATGAATAAAGAGATACAAACAATGCAAGACGTTCCATGTTCATGGGT	2092
OY	162	-----ArgTyrLeuTyrMetLeuAlaThrLeuTyrTyrLysThrTyrTyrThrAsnVal	177
		::: :	
Db	2094	AGCAGAGAACATCAATCTGCTGAATAATGGCCATCTACTGCGCCAAAGGTAAATTTAC-----	2145

Oy	177	IASpdluPheLysIle-----SerPhePheAsnLysLeuSerPhe	190
Db	2146	----AGATTCAATGCCATCCCATCAAGCTCCCAATGACTCTTCTTCCACGAAATTGGAAA	2201
Oy	190	eThrThrClyLeuPheGlyYTrPclYlIeLysArgAlaLeuLysGlnIleIleArgSerS	210
Db	2202	AACCTACTTAAAGTTCATTAATGGAACCAAAAGAGAGCCCTC-----ATTGCCAACTCATC	2255
Oy	210	nLeu--ProLeuAspIleGlyThrGlnHisSerValSerArgLeuGlnHisIleThrSer	229
Db	2256	CCTAAGCCCAAAAGACAAACCTGAGGACATCAGCTACCTGACTTCCTAAGCTTACTACAA	2315
Oy	230	SeTrLysAspTrpLysAspThrGlnIleProAlaLeuProLysPheAlaLysArgPhe	249
Db	2316	GGCTACAGTATAC-----CAAAACATCATGTGCTACTGTACCAAAA-----CAGAGATAT	2363
Oy	250	SerLeuMetValValGlnArgLeu-----AlaThr-ValAlaGlyTrpVal--	265
Db	2364	AGATCAATCGAACAAGAACAGAGCCCTCGAATAATATGCGACATCTACACATCTCAT	2423
Oy	266	----AspThrPro-----Trp-----TyrLysLysTrp--	273
Db	2424	CTTTGACCAACCTCACAACAAAACAGCAATGGGAAAGATTCCCTATTATTAATTAATGGTG	2485
Oy	274	-----TyrMetLysLeuLysAsnPheMetValAsnAr	284
Db	2484	CTGGGAAACCTGGCTACCATATGATAGAAAGCTAAACAGTATCCCTCTTACCCCTTA	2543
Oy	284	gValPheIleProThrLysLysPhePheAsnLysGlnIleArgGlnProSer--LysAl	303
Db	2544	TACAAAAATGATTCACAGATGGATTAAAGCACTTACATGTTAGCCTTAACCAATATTAAC	2603
Oy	303	aLeuLysGlnLysValSerThrAspThrLysAspLeuPheGlnAsnLysIleGlyGlnI	323
Db	2604	CCTGAGAGAAACCTAGGACCAATACCATTCAGGAC-----ATAGCATGGG	2648
Oy	323	YThrValAspPheAsnLysGlnIleArgAspProSerLysAlaLeuLysGlnLysVa	343
Db	2649	CAAG--GACTTCATGTCTTAA-----ACACAAAAGCAATGCAACAAAGC	2693
Oy	343	IserAsnAspAlaLysAspLeuPheGlnAsnLysIleGlyGlnIYThrValAspPheI	363
Db	2694	CAAAATTGACAAATGGGATCTAATTAAACTAAG-----	2722
Oy	363	eAsnAsnGlnIleArgAspProSerLysAlaLeuIleArg-----LysValSerThrCl	381
Db	2728	-----AGCTTCGCGACACAAAGAAAGTACATCAACAGTGAACAGCAACTTACAC	2786
Oy	381	YAlaGlnAspLeuPheGlnAsnLysIleGlyGlnIYThrValAspPheIleAsnAsnGl	401
Db	2781	ATGGAGAGAAATTTT-----GC	2798
Oy	401	uIleArgAspProSerLysAlaLeuIleArgLysValYTrThrGlnAlaAspAspLeuPh	421
Db	2799	AATCTACCTACATCTGACAAAGGGCTAATATTCACAGTCTCAATGAACCTCAACAAATTTA	2858
Oy	421	eGlnAsnLysIleGlyGlnIYThrValAspPheIleAsnLysGlnIleArgAspProse	441
Db	2859	CAACAAAAA-----ACAAACAACCCCATCAAAAAAGTGGGC	2894
Oy	441	rLysAlaLeuIleArgLysValSerThrGln	451
Db	2895	AAAGATATGACAGACACTTCTCCAAAGAA	2925
RESULT	25		
AAST74637			
ID	AAST74637	standard; cDNA; 11087 BP.	
XX	AAST74637:		
AC			
XX	13-FEB-2002	(first entry)	
DT			
DE		DNA encoding novel human diagnostic protein #10441.	

XX Human: chromosome mapping; gene mapping; gene therapy; forensic;  
KW food supplement; medical imaging; diagnostic; genetic disorder; ss.  
XX  
XX Homo sapiens.  
XX  
XX WO200175067-A2.  
XX  
XX PD 11-OCT-2001.  
XX  
XX PF 30-MAR-2001; 2001WO-US08631.  
XX  
XX PR 31-MAR-2000; 2000US-0540217.  
XX PR 23-AUG-2000; 2000US-0649167.  
XX  
XX PA (HXSE-) HXSEQ INC.  
XX  
XX PI Drmanac RT, Liu C, Tang YT;  
XX  
XX DR WPI: 2001-639362/73.  
XX DR P-PSDB: ABG10450.  
XX  
XX PT New isolated polynucleotide and encoded polypeptides, useful in  
PT diagnostics, forensics, gene mapping, identification of mutations  
PT responsible for genetic disorders or other traits and to assess  
PT biodiversity.  
XX  
XX PS Claim 1; SEQ ID No 10441; 103pp; English.  
XX  
XX The invention relates to isolated polynucleotide (I) and  
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,  
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome  
CC and gene mapping, and in recombinant production of (II). The  
CC polynucleotides are also used in diagnostics as expressed sequence tags  
CC for identifying expressed genes. (I) is useful in gene therapy techniques  
CC to restore normal activity of (II) or to treat disease states involving  
CC (II). (II) is useful for generating antibodies against it, detecting or  
CC quantitating a polypeptide in tissue, as molecular weight markers and as  
CC a food supplement. (II) and its binding partners are useful in medical  
CC imaging of sites expressing (II). (I) and (II) are useful for treating  
CC disorders involving aberrant protein expression or biological activity.  
CC The polypeptide and polynucleotide sequences have applications in  
CC diagnostics, forensics, gene mapping, identification of mutations  
CC responsible for genetic disorders or other traits to assess biodiversity  
CC and to produce other types of data and products dependent on DNA and  
CC amino acid sequences. AAS64197-AAS94564 represent novel human  
CC diagnostic coding sequences of the invention.  
CC Note: The sequence data for this patent did not appear in the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at ftp.wipo.int/pub/published\_pct\_sequences.  
XX  
XX SQ Sequence 11087 BP; 4422 A; 2400 C; 2076 G; 2189 T; 0 other;  
XX  
XX Alignment Scores:  
XX Pred. NO.: 2.28 Length: 11087  
XX Score: 113.00 Matches: 109  
XX Percent Similarity: 34.67% Conservative: 55  
XX Best Local Similarity: 23.04% Mismatches: 179  
XX Query Match: 4.79% Indels: 130  
XX DB: 23 Gaps: 22  
XX  
XX US-09-807-459-2 (1-458) x AAS74637 (1-11087)  
XX  
XX QY 55 LysGlySerAsnGlySerAlaSerValSerAlaTyrMetSerArgCysAlaLysGlnAsp 74  
XX DB 9544 AAGGGTATTCATTAAGCAAGAGCAAGTTCCTGCTTGGCAGACATGAT 9603  
XX  
XX QY 75 CysLeuThrLeuGln-----SerLeuLysTyrProLeuGlnAlaLysTyr--- 89  
XX DB 9604 TGTTTATCTAGAAAACCCCATCGTCTCAGCCCAAAATCTCCTTAAGCGATTAAGCAACTT 9663  
XX  
XX QY 90 --GlnProLeuThrLeuProAspPro-TyrGlnLeuGlnAlaAlaPheIleuPheLeu 108  
XX DB 10615 C-----TACTCATCTGACAAAGC 10632

DB 9664 CGGCAAGCTCAGGATACAAAATCATCATGTACAAAATCATCAAGCATCTTATACCA 9723  
QY 108 sCtLuserSpAlaAsnProAlaAsnSerThrGluLysArgPheTrpMetArgPheArgAr 128  
DB 9724 CACGACACAAACAGAGAGCCAAATCATGAGTACACCCCATTCACAAATGCTTCAAGAG 9783  
QY 128 gGlyLysAsnHisSerTyrPheHisAspLeuValPheAsnLeuLeuGluLysAsnValPh 148  
DB 9784 AATTAATA-----TACCTTAGAATCCAACTTAC 9810  
QY 148 rArgAspAlaAspAlaThrAspIleGluAsnPheAlaSer-----ArgTyrLeuTy 161  
DB 9811 AAGGATGTGAAGGACCTCTTCAGAGAACTACAAACACACTGCTCAAGAAATAAAGA 9870  
QY 162 -----ArgTyrLeuTy 165  
DB 9871 GGACACAAACAAATGGAGACATTCATGCTCATGGTAGAGAAATCAATATCGTGAA 9930  
QY 165 rMetAlaThrLeuTyrTyrTyrThrAsnValAspGluPheGlyAla----- 182  
DB 9931 AATGGCCATCTGCCCCAAGGTAATTAC-----AGATTCAATGCCATCCCCAT 9978  
QY 183 -----SerPheAsnLysLeuSerPheThrThrGlyLeuPheGlyTyrPcl 198  
DB 9979 CAAGCTACCAATGACTTCTTCACAGAAATGGAAAAAACTACTTTAAAGTTCAAAATGGA 10038  
QY 198 yLleLysArgAlaLeuLysGlnIleArgSerAsnLeu--ProLeuAspIleGlyThr 217  
DB 10039 CCAAAAGAGAGCTGC-----ATTGCCAAGTCAAATCTACAGCAAAACAAACCTGG 10092  
QY 218 GluHisSerValSerArgLeuGlnHisIleThrSerSerTyrLysAspTyrMetAspThr 237  
DB 10093 AGCGATCATGCTACCCGACTTCAAACTATACAAAGCTACAGTAT-----CAAAC 10146  
QY 238 GlnIleProAlaLeuProLysPheAlaLysArgPheSerLeuMetValGlnArgLeu 257  
DB 10147 ACACATGCTACTGCTGTAACAAA-----CAGAGTATAGACCAATGAGAACAGACAGGCC 10200  
QY 258 LeuAlaThrValAlaGlyTyr-----Val-AspThrPro----- 268  
DB 10201 CTCAGAAATTAATGCCCATATATACAACTATGATCTTCACAAACCTGACCAAAACAA 10260  
QY 269 ---Trp-----TyrLysLysTyrTyr----- 274  
DB 10261 GCAATGCGAAAGATTCCTCTTATTAATGATGCTGGAAGAAATGCTTACGCCATATG 10320  
QY 275 -----MetLysLeuLysAsnPheMetValAsnArgValPheIleProThrLysLysPh 292  
DB 10321 TAGAAAGCTGAAGACTGATCCCTTCTTACACCTTATACAAAATTAATTAAGATGAT 10380  
QY 292 ePheAsnLysGluIleArg---GluProSerLysAlaLeuLysGluLysValSerThrAs 311  
DB 10381 TAAAGACTTAATGTAGACTTAACATTAACCATTAAGAGCCATGAAGAAACCTAGGCAATAC 10440  
QY 311 pThrLysAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPheAsnLysG 331  
DB 10441 CATTCAGGC-----ATAGCATGGGCAAG---GACTTATGTCTTAA--- 10480  
QY 331 uIleArgAspProSerLysAlaLeuLysGluLysValSerAsnAspAlaLysAspLeuPh 351  
DB 10481 -----ACACCAAAAGCAATGGACCAACAAAGCAAAATTAATTAAGGATCTAAT 10530  
QY 351 eGluAsnLys-----IleGlyGlnGlyThrValAspPheIleAsnAsnGluIleAr 368  
DB 10531 TAAACCAAGAGACTCTGTCAGACAAAGAAACTTACATCAGAGTGAC---AG 10581  
QY 368 gAspProSerLysAlaLeuIleArgLysValSerThrGlyAlaGluAspLeuPheGlnAs 388  
DB 10582 GCAACCTACAAA-----TGGGAGAAATTTTCCCAAC 10614  
QY 388 nLysIleGlyGlnGlyThrValAspPheIleAsnAsnGluIleArgAspProSerLysAl 408  
DB 10615 C-----TACTCATCTGACAAAGC 10632







XX	AC	AAK68992:	
XX	DT	06-NOV-2001	(first entry)
XX	XX	Human	Immune/haematopoietic antigen genomic sequence SEQ ID NO:23804
XX	XX	Human	Immune; haematopoietic; immune/haematopoietic antigen; cancer
XX	KW	Cyclostatic; gene therapy; vaccine; metastasis; ds.	
XX	OS	Homo sapiens.	
XX	PN	MO200157182-A2.	
XX	XX	09-AUG-2001.	
XX	PD		
XX	PF	17-JAN-2001;	2001WO-US01354.
XX	XX	31-JAN-2000;	2000US-0179065.
PR	PR	04-FEB-2000;	2000US-0180628.
PR	PR	24-FEB-2000;	2000US-0184664.
PR	PR	02-MAR-2000;	2000US-0186350.
PR	PR	16-MAR-2000;	2000US-0189874.
PR	PR	17-MAR-2000;	2000US-0190076.
PR	PR	18-APR-2000;	2000US-0198123.
PR	PR	19-MAY-2000;	2000US-0205515.
PR	PR	07-JUN-2000;	2000US-0209467.
PR	PR	28-JUN-2000;	2000US-0214886.
PR	PR	30-JUN-2000;	2000US-0215135.
PR	PR	07-JUL-2000;	2000US-0216647.
PR	PR	07-JUL-2000;	2000US-0216880.
PR	PR	11-JUL-2000;	2000US-0217487.
PR	PR	11-JUL-2000;	2000US-0217496.
PR	PR	14-JUL-2000;	2000US-0218290.
PR	PR	26-JUL-2000;	2000US-0220963.
PR	PR	26-JUL-2000;	2000US-0220964.
PR	PR	14-AUG-2000;	2000US-0224518.
PR	PR	14-AUG-2000;	2000US-0224519.
PR	PR	14-AUG-2000;	2000US-0225213.
PR	PR	14-AUG-2000;	2000US-0225266.
PR	PR	14-AUG-2000;	2000US-0225267.
PR	PR	14-AUG-2000;	2000US-0225268.
PR	PR	14-AUG-2000;	2000US-0225270.
PR	PR	14-AUG-2000;	2000US-0225447.
PR	PR	14-AUG-2000;	2000US-0225757.
PR	PR	14-AUG-2000;	2000US-0225758.
PR	PR	14-AUG-2000;	2000US-0225759.
PR	PR	18-AUG-2000;	2000US-0226279.
PR	PR	22-AUG-2000;	2000US-0226681.
PR	PR	22-AUG-2000;	2000US-0226868.
PR	PR	22-AUG-2000;	2000US-0227182.
PR	PR	23-AUG-2000;	2000US-0227009.
PR	PR	30-AUG-2000;	2000US-0228924.
PR	PR	01-SEP-2000;	2000US-0229287.
PR	PR	01-SEP-2000;	2000US-0229343.
PR	PR	01-SEP-2000;	2000US-0229344.
PR	PR	01-SEP-2000;	2000US-0229345.
PR	PR	05-SEP-2000;	2000US-0229509.
PR	PR	05-SEP-2000;	2000US-0229513.
PR	PR	06-SEP-2000;	2000US-0230437.
PR	PR	06-SEP-2000;	2000US-0230438.
PR	PR	08-SEP-2000;	2000US-0231242.
PR	PR	08-SEP-2000;	2000US-0231243.
PR	PR	08-SEP-2000;	2000US-0231244.
PR	PR	08-SEP-2000;	2000US-0231413.
PR	PR	08-SEP-2000;	2000US-0231414.
PR	PR	08-SEP-2000;	2000US-0232080.
PR	PR	08-SEP-2000;	2000US-0232081.
PR	PR	12-SEP-2000;	2000US-0231968.
PR	PR	14-SEP-2000;	2000US-0232397.
PR	PR	14-SEP-2000;	2000US-0232398.
PR	PR	14-SEP-2000;	2000US-0232399.

[illegible]

Accession	Sequence	Length	Score	Percent Similarity	Best Local Similarity	Query Match	DB
US-09-807-459-2 (1-458) x AAK68992 (1-42738)							
0Y 174	TYRTThrsanValaSPGcluphneglYAlaserPhehaSNlyseurPhehThrtgLy 193	15.5	112.50	34.44%	20.39%	4.77%	22
Db 40901	TACATCATATAACACAAACGAGGAGCCAAATGATGAGTAOTCCATTCACAACTGCT	Matches: 74	Conservative: 51	Mismatches: 131	Indels: 107	Gaps: 16	
0Y 194	-----LeupheglYTPRGLYllylsarGAlaleuLysclnlelearg 208						
Db 40961	TCAAAGAGAAATAAAATACCTAGGATGCCAATTCACAGGAGTGTGAAGGCTCTTCAAG 41020						
0Y 209	serlsnleu---ProleuAsprlleglYThglnlIsseValserlrgleuglnlnlsite 227						
Db 41021	GAGAACTTCAACACCACTGTCACCAAGGAATATAAGAGAGATTCACAAACAAAGGAACATTT 41080						
0Y 228	ThlserSerlyrlylsasPrlyrmetasp---ThglnlleProAlaleuPolrysPhela 246						
Db 41081	CCATGCTCATGCGTGGTAGGAACAAATCATATGCTGAATAATGGCCATACTGCCCAAGTAATT 41140						
0Y 247	LysArgPheSerleuMetValValgln-----ArgleuLeuAla 259						
Db 41141	TATAGATTCAATGGCTTCCCATTAAGCTTACCAATGACATTTCTTCACAGAAATTGGAAAAA 41200						
0Y 260	ThlValAlaeglYrVal----- 265						

Db	41201	ACGCGTTTAACTCATATGAAACAAACAAAAAGAGCGCTGCATTGCCAACTCCAACTCAACCTAAGC	41260
Qy	265	-----	265
Db	41261	CAAAAGAACAAAGCTGGAGGCGATCACGCTACTGACTTCAACATATACTACAGGCTACA	41320
Qy	266	-----AspThrProTyrTyrIleValStryptophylMetLys	276
Db	41321	CTAACCAAAACAGCATGGTAC-----TGTATCCAAAACAGATATAGACCAATGGAAC	41374
Qy	277	-----LeuLysAsnPhenylValAsnArgValPhe	286
Db	41375	AGAACAGAGCCCTCAGAAATATACACCCACACATCTTCAAACTATCTGATCTTTGACAAACCT	41434
Qy	287	IleProthrLysLysPhe-----PheAsnLysGluIleArgGluProSer	301
Db	41435	GACAAACAAACAAATATGGGAAAGATTCCTCATTTTAATTAACGGCTGGGAAACACGG	41494
Qy	302	LysAlaLeuLysGluLysValSerThrAspThrLysAspLeuPheGluAsnLysIleGly	321
Db	41495	CTAGCCATATCTAGAACAGCTCAACCTGCATCCCTCTCTTACACCTTATATACAAAATTTAT	41554
Qy	322	GInGlyThrValAspPhePheAsnLysGluIleArgAspProSer--LysAlaLeuLys	340
Db	41555	TCAGATGGATTAAGACTTAAAC-----GTTAGACCTAAACCATATAAACCTTGAA	41608
Qy	341	GluLysValSerAsnAspAlaLysAspLeuPheGluAsnLysIleGlyGInGlyThrVal	360
Db	41609	GAAGACCTTAGCAATATCATTCAGGAC-----ATAGCGATGGGAAG---	41650
Qy	361	AspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLysValSerThr	380
Db	41651	GACTCATCTCTTAA-----ACACCAAAAGCAATGGCAACAAAGCCAAATTT	41698
Qy	381	GlyAlaGluAspLeuPheGluAsnLys-----IleGlyGInGlyThrValAspPhe	397
Db	41699	GACAAATGGATCTATTATTACTAAAGAGCTTCTGCACAGCAAAAGAAATTCACATCAGA	41758
Qy	398	IleAsnAsnGlu-----IleArgAspProSerLysAla	408
Db	41759	GCGAACAGGCACCTACAGAAATGGGAGAAAGTTTTCGATCTACTCAGCTGGACAAAGG	41818
Qy	409	LeuIleArgLysValTyrThrGluAlaAspSpleuPheGluAsnLysIleGlyGInGly	428
Db	41819	CTAAATATCCAGATCTACATGATCAACAAATAATTTCACAGAAAAAA-----	41866
Qy	429	ThrValAspPheIleAsnLysGluIleArgAspProSerLysAlaLeuIleArgLysVal	448
Db	41867	-----ACAAACAAACCCCATCAAAAAGTGGGAAAGATATGACAGACACTTC	41914
Qy	449	SerThrGlu 451	
Db	41915	TCAAAAAGAA 41923	
RESULT 28			
AC	AA35720		
XX	AA35720 standard; cDNA; 1422 BP.		
XX			
XX	AA35720;		
XX			
XX	09-JUL-1999 (first entry)		
XX			
XX	cDNA encoding a protein identified by the signal sequence trap method.		
KM	Signal sequence trap method; SST method; immunisation; inhibition;		
KM	infection; allergy; cancer; regulation; tissue formation; tissue repair		
KM	actin activity; inhibin activity; chemokine activity;		
KM	cytokine activity; blood coagulation regulation; agonist; antagonist;		
KM	metabolic disorder; hormonal disorder; immune disorder;		
KM	severe combined immunodeficiency; SCID; AIDS; thrombosis; cancer;		
XX	wound; ss.		
XX			
OS	Homo sapiens.		



AA35721,  
09-JUL-1999 (first entry)  
cDNA encoding a protein identified by the signal sequence trap method.  
Signal sequence trap method; SSR method; immunisation; inhibition;  
infection; allergy; cancer; regulation; tissue formation; tissue repair;  
activin activity; inhibin activity; chemokine activity;  
cytokine activity; blood coagulation regulation; agonist; antagonist;  
metabolic disorder; hormonal disorder; immune disorder;  
severe combined immunodeficiency; SCID; AIDS; thrombosis; cancer;  
wound; ss.  
Homo sapiens.  
WO910126-A1.  
15-APR-1999.  
06-OCT-1998; 98WO-JP04514.  
07-OCT-1997; 97JP-0274674.  
(ONOX ) ONO PHARM CO LTD.  
Fukushima D, Shibayama S, Tada H;  
WPI; 1999-277254/23.  
P-PSDB; AA02371.  
Polypeptides identified by the signal sequence trap method from a  
human cDNA library  
Claim 4; Page 141-144; 281pp; Japanese.  
AA35694-X35747 represent cDNA sequences that encode novel polypeptides  
(AA02355-84) which are identified from a human placental cDNA library  
by the signal sequence trap (SSR) method. The polypeptides have a  
broad range of physiological activity, including immunisation against  
and inhibition of infections, allergies and cancer; regulation of tissue  
formation and repair; activin/inhibin activity; chemokine/cytokine  
activity; blood coagulation regulation; and receptor/ligand agonist  
or antagonist activity. The polypeptides can be used for prevention  
and treatment of disorders including infections by bacteria, yeasts and  
viruses (including HIV) and protozoa; metabolic and hormonal disorders;  
immune disorders (including severe combined immunodeficiency (SCID)  
and AIDS; thrombosis; cancer; and traumatic or surgical wounds.  
Sequence 1613 BP; 624 A; 243 C; 359 G; 387 T; 0 other;

Alignment Scores:	
pred. No.:	0 274
Score:	111.00
Percent Similarity:	37.42%
Best Local Similarity:	20.43%
Query Match:	4.71%
DB:	20
Length:	1613
Matches:	95
Conservative:	79
Mismatches:	165
Indels:	126
Gaps:	26
	.

US-09-807-459-2 (1-458) x AAX35721 (1-1613)

Oy	62	SesalserialatyrmettsetarGcySalylslnpspcysleuthrLeugInserlu	81
	:	:   :	:::::
Dd	279	ACCAATAAATCTTATGAGAGTTTATATGATGTGCCAAC-----ACCTTGATGAGATG	332
Oy	82	LsfYlrProleuGuIualalyTyrglnProleuthreubPoaSprpyrlgInleuGl	101
	:	:     :	:   :
Dd	333	AATGCAGACTGCAGTCAAAACGTGAAGATTATTAAATGTGGATGCTTTTAAGCTCGAA	392
Oy	102	-----AlalaPhelleuPhelys	108
	:	:   :	:::
Dd	393	TCCATTAGAACCMAAAACAGACGATCATTAATGAACAGATTCCAAGATTGGAACAAGAAGAC	452

Oy	109	GlUSeRAspAlaAsnProAlaAsnSerThGluLys-----	120
Dd	453	GAAAGAACCGAATCGCTGTAGACTGGTGGAAAACCTGAAGCCTTCCTTACAAGAGAT	512
Oy	121	-----ArgPheIleTrmEArGPhenArGdlYusAnHsSerTYrPheNIsAspLeu	138
Dd	513	GTTCAAAAGTAATCAGGCATCATGAGCAAAATTGGAGTGCTATTCCAGCATTTCTTGACCAG	572
Oy	139	ValPheAsnLeuLeuGLuLvsAsnValThrArg-----	150
Dd	573	AAATTAATGGCTTCATGAGCAAAATTGCTAGACTAGACTAGACTAGAACTGAAACATAAAAA	632
Oy	151	AlaaspAlaThrAspIleGLuAsnPheAlaSerArgTYrLeuTYrMetAlaThrLeuTYr	170
Dd	633	CAGGAGACACTGCATCCATCAAGAAATATCATTGACAAACCAGAAGTAC-----	677
Oy	171	TyrIUsThrTYrAsnValAspGluPheGLyAlaSerPheAsnLysLeuSerPhe	190
Dd	678	-----TCAGTTGCAGACATTGAGCAATAAATCATGAA-----	710
Oy	191	ThrThrGLyLeuPheGLYTrrGLylleLysArgAlaLeuLysGlnlleLeArgSerAsn	210
Dd	711	-----AGAAATGAATTTGCAGCAGCACTATT--ATAAAA	740
Oy	211	LeuProLeuAspIleGLyThrGluHisSerValSerArgLeuGlnHisileThrSerSer	230
Dd	741	TTAAACCAAGACCTGGAAAGCTGACAAA-----CAGAAAGTTGGAAATGAGAGATTAAAA	794
Oy	231	Tyr-----LysAspTYrMeLAspThGlnlelPerroAlaleuProLysPheAlaLys	247
Dd	795	TATGCCAGAGCCAAAGAAAGCATTGAAACACAAATTCACAGATATCCAAATTTGGCTAG	854
Oy	248	ArgPheSerLeuMetValalGlarGLeuLeuAlaThrValAlaGLyTYrValAspThr	267
Dd	855	AAA-----TTAAACTATTTCCTTAAAGCTGCTGAGAAATTCCAAAGTTTT--	899
Oy	268	ProTPryrLysLysTPryrMetLysLeuLys-----	282
Dd	900	-----GACTTGAATTAATTAAGTTTAAATCCAGAGCTGGTGCCACTGCTCTTTC	947
Oy	283	Asn-----ArgvalPheIleProThrIlysrPhePheAsnLysGluilleArgGlu	299
Dd	948	AAATCACAGGCTCAAGTTTATGTACTCTTTAAGGAACCTCGTAATGAACTGAAGAAGAA	1000
Oy	300	ProserLYsAlaLeuLysGluLysValSer---ThrsPThrLysAsp-----	314
Dd	1008	ATTATATAAGCCCTTAATAAAAAAAAAATGGCTTGGAGGATACTTTAGACAAATTGAATGCA	1060
Oy	315	---LeupheGLuAsnLysIleGLynglLyrVal-----	325
Dd	1068	ATGATTAACAGAAAGCAAGACAGAAAGTGTGGAACTCTGAAGAAGAAAGTTCAAAAGCTGAT	1120
Oy	326	AspPhePheAsnLysGluilleArgAspProserLYsAlaLeuLysGluLysValSerAsn	345
Dd	1128	GATCTTTACCAAAAAAATTAAAGCAAGCACAGAGAAGAG--GATGAAAAATGTCCAGT	1180
Oy	346	AspAla-----LysasPheupheGLuAsnLysIleGLynglLyr	359
Dd	1185	GAGCTTGAGTCTCTTGGAAGAAACACACAGCACTGCTAGAAAGTACTGTTAAACAGGGCTC	1240
Oy	360	ValAspPheileAsnAsnGluilleArgAspProserLYsAlaLeuilleArgLysValSer	379
Dd	1245	AQTGAAGCTAG--AATGAATTTGATGCTGCTCAGCGGAATCCAAACTAGTTGTGCA	1300
Oy	380	ThrGLyalagLuAspPheGLuAsnLysIleLey-----GlngLythrValAsp	396
Dd	1302	ACCACAGCTGAAGAA-----AGACGAAAAATGGGAAATAACTTGCACACCTGCTTAGAC	1350
Oy	397	PheileAsnAsnGluilleArgAspProserLYsAlaLeu-----lleArgLysVal	413
Dd	1356	ATGCTTGCTACACATGTTGGGTCTGTAGACAAACAATCTTGAAGCAGCAGATTCGTAAGTT	1410
Oy	414	-----TyrrhgLu-----AlaaspAspLeupheGLuAsnLysIleGLynglLyr	428

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DB      1416 GATAGAGAAATATAGAGATCTCAGAGATCTCGGAAAT----- 1460
QY      429 ThrValAspPheIleuAsnLysGluIleArgAsp-----ProSerLysAlaLeuIleArg 446
DB      1461 -----ATTAAAGAGATTAGAGATAGATGAGAAAGAGCTACTCTAATT 1505
QY      447 LysValSerThrGlu 451
DB      1506 AGCTCTCTCGAGAA 1520
RESULT 30
AAAT73285
ID      AAT73285 standard; cDNA; 3278 BP.
XX
AC      AAT73285;
XX
DT      12-SEP-1997 (first entry)
DE
DE      K. lactis origin of replication complex protein 1 gene.
XX
XX      Origin of replication complex; ORC; yeast; Kluyveromyces fragilis;
XX      chromotography; peptide sequencing; primer; amplification; PCR; genome;
XX      polymerase chain reaction; open reading frame; cell growth; cancer;
XX      infection; inflammation; hypersensitivity; ds.
XX      Kluyveromyces fragilis.
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FH      Key
FT      CDS
FT      Location/Qualifiers
FT      /tag= a
FT      /product= ORC 1 protein
XX
XX      US5614618-A.
XX
XX      25-MAR-1997.
XX
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XX
XX      07-JUN-1995: 95US-0484106.
XX      16-DEC-1993: 93US-0168479.
XX
XX      (COLD-) COLD SPRING HARBOR LAB.
XX      (REGC) UNIV CALIFORNIA.
XX
XX      Bell SP, Foss M, Gavin K, Herskowitz I, Hidaka M,
XX      Kobayashi R, Laurensen P, Li J, McNally FJ, Rine J;
XX      Stillman RW;
XX
XX      WPI; 1997-201534/18.
XX      P-PSDB; AAW22230.
XX
XX      Nucleic acids encoding origin of replication complex proteins - used
XX      for screening for lead cpds. for therapy or diagnosis of disease
XX      associated with underritable cell growth
XX
XX      Claim 4: Column 59-62: 53pp: English.
XX
XX      This is the nucleotide sequence encoding the origin of replication
XX      complex protein 1 (ORC1) from the yeast Kluyveromyces fragilis. The
XX      sequence was isolated using primers based on amino acid sequence
XX      conserved between the ORC1 and SIR3 proteins from Saccharomyces
XX      cerevisiae. The amplified fragment was then used for low stringency
XX      DNA hybridisation to obtain the K. lactis ORC1 gene sequence. The ORC
XX      proteins (AAW2224-35) can be used to screen chemical libraries to
XX      identify lead compounds useful in treatment and diagnosis of undesired
XX      cell growth, e.g. cancer, infections, inflammation and hypersensitivity.
XX
SQ      Sequence 3278 BP; 1085 A; 572 C; 714 G; 902 T; 5 other:

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Alignment Scores: 0.705 Length: 3278  
Pred. No.: 111.00 Matches: 98  
Score:

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Percent Similarity: 35.50% Conservative: 82
Best Local Similarity: 19.33% Mismatches: 199
Query Match: 4.71% Indels: 128
DB: 18 Gaps: 22
US-09-807-459-2 (1-458) x AAT73285 (1-3278)
QY      5 AspSerValGlyAspValThrLysThrLeuLeuAlaIleSerGluSerValAspSerAla 24
DB      1214 GAACTATATCAGATATATGATGATCGATTTATCTGATATATCAGAAAGTAAAGAGCTTT 1273
QY      25 AlaAsnAlaIleMetIleAsnSerAsp-----MetSerAspIleLeuSerAlaValSer 42
DB      1274 GCAACCCATCTCTCTGCGACAGTGAAGAGTTTGAAGATTAACAGCTCGAGAAAG 1333
QY      43 AspAsnPheAlaGluArgIleCysSerGluVal----- 53
DB      1334 CTTCGAATTTAGACACCTGCCAGAAAGAGTCAATCTATTAAACGATATACCATTT 1393
QY      54 ---ProLysGlySerAsnGlySerAlaSerValSerAlaIleMetSer-----Arg 69
DB      1394 TCACCACTAAATATCAGACATCCATTCAGCCATCAGACAGTTCATCTCCTAGAAAG 1453
QY      70 CysAlaLysGlnAspCysLysLeuThrLeuGlnSerLeuLysThrProLeuGluAlaLysTyr 89
DB      1454 TTCCTTAAGATATATATAGTCCGCGCTAAAGAGCATATACCTATTTCCAAACGGTAT 1513
QY      90 GlnProLeuThrLeuProAspProTyrGlnLeuGluAlaPheIleLeuPheLysGlu 109
DB      1514 AAGAAATCCGAAGATTCCTGCTGACATGAT-----ATTTTCCAAAGCAT 1558
QY      110 SerAspAlaAsnProAlaAsnSerThrGluLysArgPheThrPheLysArgPheArgGly 129
DB      1559 AATAATGATTTGATATAGCGATTCATAGAGAGATTCGAACAGTTCTGCTAAAGGC 1618
QY      130 LysAsnHisSerTyrPheHisAspLeuValPheAsnLeuGluLysAsnValThrArg 149
DB      1619 AAAATGAGACATATTTTCTAAAGTCAGAGCAAGCAATTCACAGCAATAGCAAGAA 1678
QY      150 Asp---AlaAspAlaThrAspIleGluAsnPhe---AlaSerArgTyrLeuTyrMetAla 167
DB      1679 GAAATTTGCAAGCTGCTGATTTGACAAATTTATTTCCGCGAGAGAAATGATTTGCA 1738
QY      168 ThrLeuTyrTyrLysThrTyrThrAsnValAspGluPheGlyAlaSerPheAsnLys 187
DB      1739 AGTATATACCTCTCAGCTTACAGTCAATTT---GAAACGAGCACTACACCATATTAC 1795
QY      188 LeuSerPheThrThrGlyLeuPheGlyTyrPglYIleLysArgAlaLeuLysGlnIleIle 207
DB      1796 ATTGCGCGAGCGCCAGCGCTT-----GATAAACTTTGACGGTTCGAGAGTGGTT 1846
QY      208 ArgSerAsnLeuProLeuAspIleGlyThr-----GluHisSerValSerArgLeu 224
DB      1847 AAG-----GATTAATGACATCTGCACAGCAAAAGAACTTCAAGATTC 1891
QY      225 GlnHisIle-----ThrSerTyrLys----- 232
DB      1892 CAATACATTGAATCAATGCTTAAAGATTGTCAAGCAAGTATGATTAAGAGTCTTT 1951
QY      233 1952 TGGCAAAATATCTGGAGAAAGCTTACATCTGACCTGCCATGGAATCTGGAATTT 2011
QY      238 GlnIleProLeuLeuProLysPheAlaLysArgPheSerLeuMetValValGlnArgLeu 257
DB      2012 TATTTTAACAAAGTTCAGCTAGCAAAAGAGTCTGCTGCTGATTTGATGAGCTT 2071
QY      258 LeuAlaThrValAlaGlyTyrValAspThrProTyr---TyrLysLysTyr-----Tyr 274
DB      2072 GATGCAATAGTATGACAGACCAAGCATGATGATGATGATGATGATGATGATGATGAT 2131
QY      275 MetLysLeuLysAsnPheMetVal-----AsnArgValPheIleProThrLysLys 291

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Db 2132 TCAATGCGAAGCTTATGTGTAGCTGTGCGAAGACCTTAGATCTCCGAGCCAT 2191
QY 292 Phepshslnlys----- 295
Db 2192 CTGTGTAACAGATTTCGCCAGAAATGGTTTACTAGAAATATGTCATGCTTACAG 2251
QY 296 -----Gluilearg----- 298
Db 2252 CATGAGAGCTTAGACAAATCATCATTTGAGACTTAATATTTGACGAATCTGTTTC 2311
QY 299 -----GluProSerLysAlaLeuLysGluLysValSerThrAspThrLysAspLeuPhe 316
Db 2312 TATGTCGACCGGAGACGAGGTCTGACATGATCTCTCCGAGATAGTACTATATA--- 2368
QY 317 GlusnlnysIleGlyGlnGlyThrValAspPheAsnlnysGluIleArgAspProser 336
Db 2369 GAAACTGATGAGAGAAAGACGAAAGACTTCTCTAAC-----TAT 2410
QY 337 LysAlaLeuLysGluLysValSerAsnAspAlaLysAspLeuPheGluAsnLysIleGly 356
Db 2411 AAACGACTTAAACTAGATTAATCCATGATGCCATGAGATTGCATCAAGAAATTTGCT 2470
QY 357 GlnGlyThrValAspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIleArg 376
Db 2471 AGT-----GTCAGTGTGATGTGCGAGAGCTTTAAAGGTGTCAAAAGA 2515
QY 377 LysValSerThrGlyAlaGluAspLeu-----PheGluAsnLysIle--- 390
Db 2516 GCGGATGAATATGCGGAAATGATTACTTAAAGAGCTTAGATGACGACTACTCAAT 2575
QY 391 -----GlyGlnGlyThrValAspPheIleAsnAsnGluIle 402
Db 2576 TCCAAAAAAGTACTAGTGGCAATGTGACAGAAATGAAGATTAACAGAGTGTAGAAAT 2635
QY 403 ArgAspProSerLysAlaLeu 409
Db 2636 AAGCATATTACCAAGGCATTA 2656

RESULT 31
AAT62358
ID AAT62358 standard; cDNA; 3278 BP.
XX
AC AAT62358;
XX
DT 23-JUL-1997 (first entry)
XX
DE Kluyveromyces lactis origin of replication complex ORC1 gene.
XX
KW Origin of replication complex; ORC; gene therapy; cancer;
XX
neoplasia; inflammation; hypersensitivity; ds.
XX
OS Kluyveromyces lactis.
XX
FH Key Location/Qualifiers
FT misc_difference 309
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FT /note= "base 309 is given as m in the
FT misc_difference 311
FT /*tag- b
FT /note= "base 311 is given as n in the
FT misc_difference 327
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FT /note= "base 327 is given as n in the
FT misc_difference 328
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FT /note= "base 3143 is given as n in the

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FT CDS specification"
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FT /*tag- f
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PN W09640977-A1.
XX
PD 19-DEC-1996.
XX
PF 07-JUN-1996; 96WO-US09403.
XX
PR 07-JUN-1995; 95US-0484105.
XX
PA (COLD-) COLD SPRING HARBOR LAB.
PA (RESC ) UNIV CALIFORNIA.
PI Bell SP, Foss M, Herskowitz I, Kobayashi R, Laurensen P;
PI Li J, McNally FJ, Rine J, Stillman BW;
PI
DR MPI; 1997-052354/05.
DR P-PSDB; AAM14136.
XX
PT Nucleic acid encoding origin of replication complex (ORC) protein -
PT useful to screen for lead pharmaceuticals capable of disrupting ORC
PT protein function, and inhibiting cell growth
XX
PS Disclosure: Page 16-18; 57pp; English.
XX
CC Isolated cDNA clones (AAT62358-63) respectively encode origin of
CC replication (ORC) proteins (AAM14136-41) from Kluyveromyces lactis,
CC Schizosaccharomyces pombe, human (ORC1), Arabidopsis thaliana,
CC Caenorhabditis elegans and human (ORC2). The K. lactis ORC1 clone
CC was obtained by PCR amplification using primers based on Saccharomyces
CC cerevisiae ORC1 and SIR3 sequences. The isolated nucleic acids can
CC be utilised in the prodn. of ORC polypeptides, to design probes and
CC primers for the detection and amplification of ORC genes, and in
CC gene therapy appls., e.g. antisense oligonucleotides capable of
CC inhibiting the intracellular expression of a targeted ORC
CC transcript.
XX
SQ Sequence 3278 BP; 1085 A; 572 C; 714 G; 902 T; 5 other:

Alignment Scores:
Pred. No.: 0.705 Length: 3278
Score: 111.00 Matches: 98
Percent Similarity: 35.50% Conservative: 82
Best Local Similarity: 19.33% Mismatches: 199
Query Match: 4.71% Indels: 128
DB: 18 Gaps: 22

US-09-807-459-2 (1-458) x AAT62358 (1-3278)
QY 5 AspSerValGlyAspValThrIysThrLeuLeuAlaIleSerGluSerValAspSerAla 24
DB 1214 GAAGCTATATCAGATAATGCAATCGATTATATGCAATATACGAAAGTAAAGAGATT 1273
QY 25 AlaAsnAlaIysrMetIleAsnSerAsp-----MetSerAspTyrLeuSerAlaValser 42
DB 1274 GCAAAAGCATCTCTTCGACAGATGATGAAGAGTTTGAAGATTTACAGCTCTGCAGAAAG 1333
QY 43 AspAsnPheAlaGluArgIleCysSerGlnAla----- 53
DB 1334 CTTCGAATGTGTAACCTCCCAAGAAAGTGCAGATCTATTAAACCATATATACCAT 1393
QY 54 ---ProLysGlySerAsnCysSerAlaSerValSerAlaTyrMetSer-----Arg 69
DB 1394 TCACCAAGTAAATACACAGATCCATTTGCAGCCATGCGAGTTTCATTCATCTCCTAGAAAG 1453
QY 70 CysAlaLysGluAspCysLeuThrLeuGlnSerLeuLysTyrProLeuAlaLysTyr 89
DB 1454 TTCCTTAAAGATATATATATGCGCGCTAAAGAGCATATATCTCTTCCAAAGCGTAT 1513
QY 90 GlnProLeuThrLeuProAspProTyrGlnLeuGluAlaAlaPheIleLeuPheLysGlu 109
DB 1513

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Db 1514 AAGAAATCGAAGATTCCTGACTTGACGAT-----ATTTCACAAAGCAT 1558
OY 110 SerAspAlaSerProAlaSerThrGluLysArgPheThrMetArgPheArgGly 129
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 1559 AATATATATTTGGATATAGCTGCAATTAGAGAGGATTCAGACAGCTTTGCTTAAAGGC 1618
OY 130 LysAsnHisSerThrPheHisAspLeuValPheAsnLeuGluLysAsnValThrArg 149
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 1619 AATATGAGACTATTTTCTTAAGCTGACAGAACGAAATTCGAAAGAAATGAGAAAGAA 1678
OY 150 Asp---AlaAspAlaThrAspIleGluAsnPhe---AlaSerArgThrLeuThrMetAla 167
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 1679 GAAATGTCGAAGCTGCTGCAATTCGACAAATATCTCCGCGAAGAAATGAAATTTGCA 1738
OY 168 ThrLeuThrThrLysThrThrAsnValAspLupheGluAlaSerPhePheAsnLys 187
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 1739 AGTATATACCTCTTCACCTTACAGTCAATT---GACAGAGCACTAGACACCAATTTTAC 1795
OY 188 LeuSerPheThrThrGlyLeuPheGlyTrpGlyLysArgAlaLeuLysGluIleIle 207
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 1796 ATTGCCGGAGCGCCAGCGCTT-----GCTAAACTTTGACGGTTCCGAGAGTGGTT 1846
OY 208 ArgSerAsnLeuProLeuAspIleGlyThr-----GluHisSerValSerArgLeu 224
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 1847 AAG-----GATTTAATGACATCTGCAGACCAAAAGAACTTCCAAAGATTTC 1891
OY 225 GluHisIle-----TherSerThrLys----- 232
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 1892 CAATACATTGAAATCATGCTTAAAGATTGTCAAAACAGATGATAGTATGAAAGTCTTT 1951
OY 233 -----AspTyrMetAspThr 237
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 1952 TGGCAAAATATCTGAGAAAGCTTACATCTGAGCTGCATGCAATCTCGAGTTT 2011
OY 238 GluIleProAlaLeuProLysPheAlaLysArgPheSerLeuMetValValGluArgLeu 257
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 2012 TATTTTAAACAAGATTCAGCTACGAAAAACGTCCTATCGTGTGTTAGTAGAGCTT 2071
OY 258 LeuAlaThrValAlaGlyTyrValAspThrProTrp---TyrLysLysTrp---Tyr 274
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 2072 GATGCAATTAGTACAGAGCAAGATGATATGACAACTCTTAAGTGGCGTACCTAT 2131
OY 275 MetLysLeuLysAsnPheMetVal-----AsnArgValPheIleProThrLysLys 291
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 2132 TCAAAATCGAAACTTATGTGTAGCTGTGCAAAACACTTATGATCTCCGACAGCCAT 2191
OY 292 PhePheAsnLys----- 295
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 2192 CTTCGTAAACAAGATTCCTCAGAAATTCGTTTACTAGAAATTATGTTCACTGTTACACG 2251
OY 296 -----GluIleArg----- 298
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 2252 CATGAGAGCGCTTAGAACATCATCATTTGAGACTTAATTTGACGAATCTAGTTTC 2311
OY 299 -----GluProSerLysAlaLeuLysGluLysValSerThrAspThrLysAspLeuPhe 316
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 2312 TATGTCGACCCGGAGACAGGAGTTCGTACATGATCTCTCGGATAGTACTATATA--- 2368
OY 317 GluAsnLysIleGlyGlnGlyThrValAspPhePheAsnLysGluIleArgAspProSer 336
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 2369 GAAACATCATGACAGAAAGAAAGCGAAAGACTTCTCTAAC-----TAT 2410
OY 337 LysAlaLeuLysGluLysValSerAsnAspAlaLysAspLeuPheGluAsnLysIleGly 356
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 2411 AAACGACTAAACCTTAAATCTGATGCTCATGATTCATCAAGAAATATGCT 2470
OY 357 GlnGlyThrValAspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIleArg 376
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 2471 AGT-----GTCAGGTGTGATGTGCGAGAGCTTTAAAGGTGTGCTAAAGA 2515
OY 377 LysValSerThrGlyAlaGluAspLeu-----PheGluAsnLysIle--- 390
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 2516 GCGGTACAAATATGCGAAATGATTACTTAAAGAGCTTACATATGACCGCATGTCAAT 2575

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OY 391 -----GlyGlnGlyThrValAspPheIleAsnGluIle 402
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 2576 TCCAAAAAAGTACTAGTGGCAATGCTACAGGAATGACAAATTACAGAGTGAATTT 2635
OY 403 ArgAspProSerLysAlaLeu 409
    : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 2636 AACCATATTACCAAGCATTTA 2656

RESULT 32
AAQ06842
ID AAQ06842 standard; RNA; 1777 BP.
XX
AC AAQ06842;
XX
XX 05-MAR-1991 (first entry)
DT
XX RNA segment 4 coding for HA.
DE
XX Influenza; haemagglutinin; HA; ribosomal frameshift signal sequence;
KM membrane anchor; RFS; ss.
XX
OS Influenza virus A/PR8/34.
XX
FH Key Location/Qualifiers
FT CDS 33..1732
FT sig_peptide 33..82
FT /*tag- b 83..1063
FT mat_peptide /*tag- c
FT /*product- HA1 1064..1730
FT /*tag- d
FT mat_peptide /*product- HA2 1632..1692
FT /*tag- e
FT /*label- insertion site for RFS

WO9014422-A.
XX
XX 29-NOV-1990.
XX
XX 21-MAY-1990; 90WO-G800791.
XX
XX 19-MAY-1989; 89GB-0011555.
XX
PA (LYNX-) LYNXVALE LTD.
XX
XX Inglis SC, Brierley I;
XX
XX WPI; 1990-375989/50.
XX
PT Ribosomal frame shifting signal sequences - isolated from
PT infectious bronchitis virus genomic RNA and used in protein
PT prodn.
XX
XX Disclosure; Fig 19; 55pp; English.
XX

The HA gene encodes a spike-like protein which is embedded in the
membrane via a hydrophobic anchor sequence. A portion of this
anchor sequence may be replaced with a ribosomal frame shift signal
sequence (RFS), in such a way that ribosomes translating the new
HA sequence will usually terminate before the hydrophobic sequence
is encountered, leading to the prodn. of a secreted form of the HA.
It has been found that the primary sequence of the RFS can be rad-
ically altered as long as the the secondary and tertiary structures
are preserved, so it is possible to design an RFS which encodes
CC hydrophobic amino acids, and therefore preserves the integrity of
CC the anchor.
CC See also AAQ06841 and AAQ07007.
CC
XX Sequence 1777 BP; 621 A; 331 C; 408 G; 417 U; 0 other;
SQ

```



## Alignment Scores:

Pred. NO.: 0.391 Length: 1777  
 Score: 110.00 Matches: 97  
 Percent Similarity: 35.04% Conservative: 67  
 Best Local Similarity: 20.73% Mismatches: 170  
 Query Match: 4.66% Indels: 134  
 DB: 11 Gaps: 24

US-09-807-459-2 (1-458) x AA006842 (1-1777)

```

OY 23 SerAlaAlaSerAlaTyrMetIleAsnSerAspMetSerAspTyr----- 37
DB 332 AACUCUGAAGUAGAAUUGUUUACAGGAGAUUUUACUAGCAGUAGAGAGAGGAG 391
OY 38 ---LeuSerAlaValSerAspAsnPhenAlaGluArgIleCysSerGlnValProIleGly 56
DB 392 CAUUGGAGCUCAGUGUCA-----UUCGAAAGAUUC---GAAAUUUUUCCAAAGAA 442
OY 57 Ser-----AsnCysSerAlaSerValSerAlaTyrMetSerArgCysAla 71
DB 443 AGCUCUAGGCCCAACACACACAAAGAGUAGCGGCAUUGCUCCCAUGCGGGG 502
OY 72 LysGlnAsp-----CysIleuThrLeuGlnSerLeuLysTyrProIleu 85
DB 503 AAAACGAGUUUUUACGAAAUUUUGCUAUGGCGAGAGAGAGGCGCUAACCACCA--- 559
OY 86 GluAlaLysTyrGlnProIleuThrLeuProAspProTyrGlnLeuGlnAlaIaPhelle 105
DB 560 -----AACGUGAAAAUUUUUUUUGG 580
OY 106 LeuPheLysGluSerAspAla-----AspProAlaAsnSerThr 118
DB 581 AACAGAAAGGAAAGAAAGUCCUUUACUGUGGGUUUACAUACACCGGCUAACAGUAG 640
OY 119 GluLysArgPheTyrMetArgPheArgGlyLysAsnHisSerTyrPheHisAspLeu 138
DB 641 GAUCCAACAG-----AAUUAUCUACAGAUAGAAAU---GCUUAUGUCUCUGUAGUG 688
OY 139 ValPheAsnLeuLeuGlnLysAsnValThrArgAspAlaAspAlaThrAspIleGluAsn 158
DB 689 ACUUCGAAAUUUUACAGGAGAUUUUACCCCGGAAUAGCGAAAGAACCCAAAUUAGAGAGU 748
OY 159 PheAlaSerArgTyrLeuTyrMetAlaThrLeuTyrTyrTyrTyrThrAsnValAsp 178
DB 749 CAAGCUGGAGAGAUACUUAUACUGGACCUUGCUAAAAACCGGAGACACAAUUAUUUU 808
OY 179 GluPheGlyAlaSerPhePhe-----AsnLysLeuSerPheThrThrGlyLeuPhe 195
DB 809 GAGCGCAAUUGAAUUCUUAUACGACCAAGUUAUGUUUGCAGUCAGUAGAGGC---UUU 865
OY 196 GlyTyrPglYIleLysArgAlaLeuLysGlnIleIleArgSerAsnLeuProLeuAspIle 215
DB 866 GGGUCCGGC-----AUCACACCCUCAAAGCAAGCAUCAAUAGUAG 904
OY 216 GlyThrGlnHisSerValSerArgLeuGlnHisIleThrSerSer-----TyrLysAsp 233
DB 905 UGUAAACAGAGAGUUGCAACACCCUUGGAGAGUUAUAAACAGCAGUCUCCUUUCCAGAU 964
OY 234 TyrMetAspThrGlnIleProAlaLeuProLysPheAlaLysArgPheSerLeuMetVal 253
DB 965 AUAACACCGAGUCACAUAUUGAGAGUGCCAAAUUAGUUCAGAGGCCAAUUGAGAGUG 1024
OY 254 Val-----GlnArgLeuLeuAlaThrValAla 262
DB 1025 GUUACAGAGACUAAAGACAUUCCGCUUACUUAUCCAGGCGUUAUUUGAGCCAUUCC 1084
OY 263 GlyTyrValAspThrProTyr-----TyrLysTyrTyrTyr----- 274
DB 1085 GGUUUUUUUAAGAGGGAGGAGCUGAGAUAGAUAGAGUGGUGUUAUUCUACUACUGAG 1144
OY 275 -----MetLysLeuLysAsnPh 280
  
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DB 1145 AAUAGACAGGAGUACAGGCUUACGACGCGAUCAAAAAAGCACACAAAUAUCCAUUAAACGG 1204
OY 281 MetValAsnArgVal-----PheIleProThrLys 290
DB 1205 AUUACAAACAGAGUGUAAUCUUGUAUCGAGAAAUACAAUUCAUUCCACAGCUGUGGU 1264
OY 291 LysPhePheAsnLysGluIleArgGluProSerLysAlaLeuLysGluLysValSerThr 310
DB 1265 AAAGAUAUCCAAACAA---UUAGAAAAAGAGUAGAAAUUUUAUUAUAAAAAGUUGAU 1321
OY 311 AspThrLysAsp-----LeuPheGlnAsnLysIle 320
DB 1322 GGAUUUUCUGACAUUUGGACAUUAUACGAGAUUGCUUAGUUCUACUUGGAAAAU----- 1375
OY 321 GlyGlnGlyThrValAspPheAsnLysGluIleArgAspProSerLysAlaLeuLys 340
DB 1376 ---GAAAGACUCCUGAUUUUCCAUAGACUCAAUUGUCAAAUCCUGAUAGAAAGUAAAA 1432
OY 341 GluLysValSerAsnAspAlaLysAspLeuPheGluAsnLysIleGlyGlnGlyThrVal 360
DB 1433 AGCCAAUUAUAAAGAUUAUAGUCCAAAGAA-----AUCGAAUUGGAGUUGUU 1477
OY 361 AspPhe-----IleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLys 377
DB 1478 GAGUUCUACACAGUUGUACAAUGAA-----UCCAUUGGAAAGU 1516
OY 378 ValSerThrGlyAlaGluAspLeu-----PheGluAsnLysIleGlyGlnGly 393
DB 1517 GUUAGAAUUGGACUUAUAGAUUCCCAAAUUAUUCAGAAAGAGUCAAAGUUAACAGGAA 1576
OY 394 ThrValAspPheIleAsnAsnGlu 401
DB 1577 AAGGUAGAGUAGUAGUAAUUGGAA 1600

RESULT 33
AA574418
ID AA574418 standard; cDNA; 1816 BP.
XX
AC AA574418;
XX
DE 13-FEB-2002 (first entry)
XX
DE DNA encoding novel human diagnostic protein #10222.
XX
KM Human; chromosome mapping; gene mapping; gene therapy; forensic;
XX food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
OS Homo sapiens.
XX
PN W0200175067-A2.
XX
PD 11-OCT-2001.
XX
PF 30-MAR-2001; 2001WO-0508631.
XX
PR 31-MAR-2000; 2000US-0540217.
XX
PR 23-AUG-2000; 2000US-0649167.
XX
PA (HYSE-) HYSEQ INC.
XX
PI Drmanac RT, Liu C, Tang YT;
XX
DR WPI: 2001-639362/73.
XX
DR P-PSDB: ABG10231.
XX
PT New isolated polynucleotide and encoded polypeptides, useful in
XX diagnostics, forensics, gene mapping, identification of mutations
XX responsible for genetic disorders or other traits and to assess
XX biodiversity -
XX
PS Claim 1; SEQ ID NO 10222; 103pp; English.
XX
CC The invention relates to isolated polynucleotide (I) and
  
```

CC polypeptide (II) sequences. (I) is useful as hybridisation probes,  
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome  
CC and gene mapping, and in recombinant production of (II). The  
CC polynucleotides are also used in diagnostics as expressed sequence tags  
CC for identifying expressed genes. (I) is useful in gene therapy techniques  
CC to restore normal activity of (II) or to treat disease states involving  
CC (II). (II) is useful for generating antibodies against it, detecting or  
CC quantitating a polypeptide in tissue, as molecular weight markers and as  
CC a food supplement. (II) and its binding partners are useful in medical  
CC imaging of sites expressing (II). (I) and (II) are useful for treating  
CC disorders involving aberrant protein expression or biological activity.  
CC The polypeptide and polynucleotide sequences have applications in  
CC diagnostics, forensics, gene mapping, identification of mutations in  
CC responsible for genetic disorders or other traits to assess biodiversity  
CC and to produce other types of data and products dependent on DNA and  
CC amino acid sequences. AAS64197-AAS94564 represent novel human  
CC diagnostic coding sequences of the invention.  
CC Note: The sequence data for this patent did not appear in the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at [ftp.wipo.int/pub/published\\_pct\\_sequences](http://wipo.int/pub/published_pct_sequences).

XX Sequence 1816 BP; 726 A; 380 C; 328 G; 382 T; 0 other;

Alignment Scores:  
Pred. NO.: 0.45 Length: 1816  
Score: 109.50 Matches: 76  
Percent Similarity: 37.99% Conservative: 41  
Best Local Similarity: 24.68% Mismatches: 113  
Query Match: 4.64% Indels: 78  
DB: 23 Gaps: 16

US-09-807-459-2 (1-458) x AAS74418 (1-1816)

QY 183 serPhepHeaNsLysLeuSerPheThrGlyLeuPheGlyTyrPglYlleYsArgLa 202  
:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||  
Db 721 ACTTCTTCACAGAAATTCGAAATTCCTTAAGTTCATCAATGACCAACCAAAAGACC 780  
QY 203 LeuYsGlnIlelleArgSerAsnLeu--ProLeuAspIleGlyThrGlnHisSerValS 222  
|||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||  
Db 781 TGC-----ATTGCCAAGTCAATCTTAAGCCAAAGAACGAGCATCATCTGA 834  
QY 222 eArYgLeuGlnHisIleThrSerSerYrYsAspTyrMeLAspThrInleProAlaL 242  
:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||  
Db 835 CCCGACTTCAACACTATCTATACAGGCTACAGTAAAT-----CAAAACAGCATGTGACTGG 888  
QY 242 euProYsPheAlaLysArPheSerLeuMetValIValGlnArgLeuDeuAlaThrValA 262  
:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||  
Db 889 TACCAAA-----CAGGATATAGACCAATGGAACAGACAGACCCTCAGAAATATG 942  
QY 262 laGlyTyr-----Val-AspThrPro-----Tyr----- 269  
|||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||  
Db 943 CCGCATATCTACAACTATCTGATCTTCGACAAACCTGACCAAAACAGCATGGGGAAG 1002  
QY 270 -----TyrLysLysTyrTyr-----MetLys 276  
:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||  
Db 1003 GATTCCTATTATTAATGGTACTGCGGAAACCTGACCATATGTAAGAGCTGAAA 1062  
QY 277 LeuYsAsnPhMeTValAsnArgValPheIleProThrLysLysPheAsnLysGlu 296  
|||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||  
Db 1063 CTAATCCCTTCCTTACACCTTATACAAAAATTAATTAACATGATTAAGACTTAAT 1122  
QY 297 IleArg--GluProSerLysAlaLeuYsGluYsValSerThrAspThrLysAsnLeu 315  
:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||  
Db 1123 GTTAGACTTAACCAATTAAGAGCCCTAGACAAACCTAGCAATACATTCAGAGAC--- 1179  
QY 316 PheLysAsnLysIleGlyGlnGlyThrValAspPhePheAsnLysGluIleArgAspPro 335  
:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||  
Db 1180 -----ATAGGCAATGGCAAG--GACTTCATGTCTTAA-----ACA 1212  
QY 336 SerLysAlaLeuYsGluYsValSerAsnAspAlaLysAspLeuPheGlnAsnLys--- 354  
|||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||  
Db 1213 CCAAAAGCAATGGCAACAAAGACAAATTCACAAATGGGATCTAATTAACCAAGAGGC 1272

QY 355 -----IleGlyGlnGlyThrValAspPheIleAsnGlnIleIleArgAspProSerLys 372  
:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||  
Db 1273 TTCTGCACAGCAAAAGAAACATACATCATGAGGTGAAC-----AGGCAACCTTCAAAA 1323  
QY 373 AlaLeuIleArgLysValSerThrGlyAlaGlnAspLeuPheGlnAsnLysIleGlyGln 392  
|||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||  
Db 1324 -----TGGCAGAAATTTTCGCAAC----- 1344  
QY 393 GlyThrValAspPheIleAsnGlnIleIleArgAspProSerLysAlaLeuIleArgLys 412  
1345 -----TACTCATCTGCACAAAGGCTTAATTCACGA 1374  
QY 413 ValYrThrGlnAlaAspAspLeuPheGlnAsnLysIleGlyGlnGlyThrValAspPhe 432  
|||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||  
Db 1375 AACTCATGTGAACCTCAACCAATTTACAGAAAGAAAA-----ACAATGGGCCCC 1422  
QY 433 IleAsnLysGlnIleArgAspProSerLysAlaLeuIleArgLysValSerThrGlnAla 452  
|||:::|||||:::|||||:::|||||:::|||||:::|||||:::|||||  
Db 1423 ATCAAAAGGTGGCGAGAGATATGAAACAGCATCTTCAAAAGAACACATATGACGCC 1482  
QY 453 AspaSnLeuGlnLys 458  
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Db 1483 AAAAAACATGAAAAAA 1500

RESULT 34  
AAS69641  
ID AAS69641 standard; cDNA; 2277 BP.

XX AAS69641;

DT 13-FEB-2002 (first entry)

DE DNA encoding novel human diagnostic protein #5445.

KW Human: chromosome mapping; gene mapping; gene therapy; forensic;  
KW food supplement; medical imaging; diagnostic; genetic disorder; ss.

OS Homo sapiens.

PN WO200175067-A2.

XX 11-OCT-2001.

PD 30-MAR-2001; 2001WO-US08631.

PF 31-MAR-2000; 2000US-0540217.

PR 23-AUG-2000; 2000US-0649167.

XX (HYSE-) HYSEQ INC.

PI Drmanac RT, Liu C, Tang YT;

DR WPI: 2001-639362/73.

XX P-PSDB; ABG05454.

PT New isolated polynucleotide and encoded polypeptides, useful in  
PT diagnostics, forensics, gene mapping, identification of mutations  
PT responsible for genetic disorders or other traits and to assess  
PT biodiversity

PS Claim 1; SEQ ID No 5445; 103pp; English.

XX The invention relates to isolated polynucleotide (I) and  
XX polypeptide (II) sequences. (I) is useful as hybridisation probes,  
XX polymerase chain reaction (PCR) primers, oligomers, and for chromosome  
XX and gene mapping, and in recombinant production of (II). The  
XX polynucleotides are also used in diagnostics as expressed sequence tags  
XX for identifying expressed genes. (I) is useful in gene therapy techniques  
XX to restore normal activity of (II) or to treat disease states involving  
XX (II). (II) is useful for generating antibodies against it, detecting or  
XX quantitating a polypeptide in tissue, as molecular weight markers and as  
XX a food supplement. (II) and its binding partners are useful in medical  
XX imaging of sites expressing (II). (I) and (II) are useful for treating



Pred. No.:	0.609	length:	2277
Score:	109.50	Matches:	73
Percent Similarity:	37.21%	Conservative:	39
Best Local Similarity:	24.25%	Mismatches:	107
Query Match:	4.64%	Indels:	82
DB:	23	Gaps:	15

US-09-807-459-2 (1-458) x AAS71279 (1-2277)

OY	183	SerPheAsnLysLeuSerPheThrGlyLeuPheGlyTyrPoliLysArgAla	202
Db	868	ACTTCTTCCACGAAATTGGAAAAACTACTTTAAAGTCTCATGTGAACCAAAAAAGACC	927
OY	203	LeuLysGlnLeileIleArgSerAsnLeu--ProLeuAspIleGlyThrGlnHisSerValS	222
Db	928	CTC-----ATCGCCAAAGTCATATCTGTGACCAAAAGACAAAGCCGAGAGCATTCAGCTA	981
OY	222	erAlaGlyLeuGlnHisIleThrSerSerTyrLysAspIlyrMetAspThrGlnIleProAlaI	242
Db	982	CTGTACTCTCGAATCATATCTACAAAGCCTACAGTAAAC-----CAAAACACCCTGGTACTCG	1035
OY	242	euprOlysPheAlaLysArgPheSerLeuMetValAlaGlnArgLeuLeuAlaThrValA	262
Db	1036	TACAAAA-----CAGAGATATAGACCAATGGAACAGAAAGAGCCCTCAGAAATATAG	1089
OY	262	laGlyTyr-----Val-AspThrPro-----	268
Db	1090	CCGCTATCTACAACTATCTGATCTCTTGACAAACCTCACAAAACAGCAATGGCGAAG	1149
OY	269	-----TyrLysLysTyrTyrMetLysLysAsnPheMet	281
Db	1150	GATTCCTCATTTTAAATATGTCGTGGGAAAAATGGCTAACCTATGTGAAAGACTGAA	1205
OY	282	ValAsnArgValPheIleProThrLysLysPheAsnLysGluIleArgLys-----	299
Db	1210	CTGTGATCCCTCTCCCTTACACCTTATACAAAATTAATTCAGATGATTAACACTTACAT	1265
OY	300	-----ProSerLysAlaLeuLysGluLysValSerThrAspThrLysAspLeu	315
Db	1270	GTTAGAGCTAAACCATTAATAACCTTGAAGAAAACTAGGCATATCCATTCAAGGAC--	1322
OY	316	PheGluAsnLysIleGlyGlnGlyThrValAspPhePheAsnLysGluIleArgAspPro	335
Db	1327	-----ATAGGCATGGCGAAG--GACTTCATGCTTAA-----CCA	1355
OY	336	SerLysAlaLeuLysGluLysValSerAsnAspAlaLysAspLeuPheGluAsnLys--	354
Db	1360	CCAAAGCAACGCGACGAAAGCCAAATTGACAAATGGAGTCTAATTAACTAAAGAAC	1419
OY	355	-----IleGlyGlnGlyThrValAspPheIleAsnAsnGluIleLeuArgAspProSerLys	372
Db	1420	TTCGTGCACAGCAAAAGAACTACCATCAGAGTAAAC-----AGGCACCTCAAA	1477
OY	373	AlaLeuIleArgLysValSerThrGlyAlaGluAspLeuPheGluAsnLysIleGlyGln	392
Db	1471	-----TGGGAGAAATTTTCACAAAC-----	1499
OY	393	GlyThrValAspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLys	412
Db	1492	-----TACTCATCTGACCAAAAGCGCTAATATCCAA	1521
OY	413	ValTyrThrGluAlaAspAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPhe	432
Db	1522	ATCTACATGAATGACCTCCGCAAAATTTTACAGAAAAA-----	1557
OY	433	IleAsnLysGluIleArgAspProSerLysAlaLeuIleArgLysValSerThrGlu	451
Db	1558	ACAACAACCAACCAACCAAAAGTGGCGAAGACATGAACAGACACTTCTCAAAAGAA	1614
RESULT	36		
AA574306			
ID	AA574306	standard; cdna; 2277 bp.	

AC	AA574306;
XX	
DT	13-FEB-2002 (first entry)
XX	
DE	DNA encoding novel human diagnostic protein #10110

KW Human; chromosome mapping; gene mapping; gene therapy; forensic;  
KW food supplement; medical imaging; diagnostic; genetic disorder; ss

OS	Homo sapiens.
XX	
PN	WO200175067-A2.

PD	11-OCT-2001.
XX	
PF	30-MAR-2001; 2001WO-US08631
XX	
PR	31-MAR-2000; 2000US-0540217
PR	23-AUG-2000; 2000US-0649167

PA (HYSE-) HYSEQ INC.

PI Drmanac RT, Liu C, Tang YT;

DR WPI; 2001-639362/73.

DR P-PSDB; ABG10119.

PT New isolated polynucleotide and encoded polypeptides, useful in diagnostics, forensics, gene mapping, identification of mutations responsible for genetic disorders or other traits and to assess biodiversity -

PS Claim 1; SEQ ID No 10110; 103pp; English.

CC The invention relates to isolated polynucleotide (I) and  
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,  
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome  
CC and gene mapping, and in recombinant production of (II). The  
CC polynucleotides are also used in diagnostics as expressed sequence tags  
CC for identifying expressed genes. (II) is useful in gene therapy techniques  
CC to restore normal activity of (I) or to treat disease states involving  
CC (II). (II) is useful for generating antibodies against it, detecting or  
CC quantitating a polypeptide in tissue, as molecular weight markers and as  
CC a food supplement. (II) and its binding partners are useful in medical  
CC imaging of sites expressing (II). (I) and (II) are useful for treating  
CC disorders involving aberrant protein expression or biological activity.  
CC The polypeptide and polynucleotide sequences have applications in  
CC diagnostics, forensics, gene mapping, identification of mutations  
CC responsible for genetic disorders or other traits to assess biodiversity  
CC and to produce other types of data and products dependent on DNA and  
CC amino acid sequences. A564197-A594564 represent novel human  
CC diagnostic coding sequences of the invention.  
CC Note: The sequence data for this patent did not appear in the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at [ftp.wipo.int/pub/published\\_pat\\_sequences](http://wipo.int/pub/published_pat_sequences).

**sq** Sequence 2277 BP; 878 A; 483 C; 473 G; 443 T; 0 other;

Alignment Scores:	
Pred. No.:	0.609
Score:	109.50
Percent Similarity:	37.21%
Best Local Similarity:	24.25%
Query Match:	4.64%
DB:	23
Length:	2277
Matches:	73
Conservative:	79
Mismatches:	107
Indels:	82
Gaps:	15

US-09-807-459-2 (1-458) x AAS74306 (1-2277)

Qy	183	SerPhepheanbLysLeuSerPhehLrThGcLyeuDehGeyTrpGcLyIeLysaGAla	202
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Db	868	ACTTTCTTCCAGAAATTGGAAAAAACTACTTTTAAAGTTCAATATGGAAACCAAAAAAGACCC	927
Oy	203	LeuLysGlnIleIleLysSerAsnLeu--ProLeuAspIleGlyTrhGlnHisSerValS	222



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QY 269 -----TTP---TyrLysLysTrpTyrMetLysLysAsnPhmet 281
DB 1150 GATTCCCTATTATATAATGCTGCGGAAAAAATGCTAACCAATATGTAAGAAACCTGAAA 1209
QY 282 ValAsnArgValPheIleProThrLysLysPheAsnLysGluIleArgGlu----- 299
DB 1210 CTGATCCCTCTTCCCTTACCACTTATACAAAATTAATTCAGATGATGATTAAGACTTACAT 1269
QY 300 -----ProSerLysAlaLeuLysGluLysValSerThrAspThrLysAspLeu 315
DB 1270 GTTAGAGCTAAACCACTAAAACCTTAGAAGAAAAGAACTAGCAATACCATTCAGAC--- 1336
QY 316 PheGluAsnLysIleGlyGlnGlyThrValAspPhePheAsnLysGluIleArgAspPro 335
DB 1327 -----ATAGCATGGGCAAG---GACTTCATGCTTAA-----CCA 1359
QY 336 SerLysAlaLeuLysGluLysValSerAsnAspAlaLysAspLeuPheGluAsnLys--- 354
DB 1360 CCAAAACCAACGGCAGCAAGCAAAATTCACAAATGGATCTTAATTAACCTAAAGAAC 1419
QY 355 -----IleGlyGlnGlyThrValAspPheIleAsnAsnGluIleArgAspProSerLys 372
DB 1420 TTCTGCACAGCAAAAGAACTACCATCAGACTGAC-----AGCAACCTACAAA 1470
QY 373 AlaLeuIleArgLysValSerThrGlyAlaGluAspLeuPheGluAsnLysIleGlyGln 392
DB 1471 -----TGGGAGAAATTTTCACACAC----- 1491
QY 393 GlyThrValAspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLys 412
DB 1492 -----TACTCATCTGACAAAGGCTTAATTCACCA 1521
QY 413 ValTyrThrGluAlaAspAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPhe 432
DB 1522 ATCTACATGACATCCGCAACATTTACAGAAAA----- 1557
QY 433 IleAsnLysGluIleArgAspProSerLysAlaLeuIleArgLysValSerThrGlu 451
DB 1558 ACAAAACCAACCCCAACAAAGTGCGCAGACATGAACACACTCTCAAAAGAA 1614
RESULT 38
AAS74599
ID AAS74599 standard; cDNA; 2277 BP.
XX
AC AAS74599;
XX
DT 13-FEB-2002 (first entry)
XX
DE DNA encoding novel human diagnostic protein #10403.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
OS Homo sapiens.
XX
WO200175067-A2.
XX
PD 11-OCT-2001.
XX
PF 30-MAR-2001; 2001MO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.
PR 23-AUG-2000; 2000US-0649167.
XX
PA (HYSE-) HYSEQ INC.
PI Drmanac RT, Liu C, Tang YT.
XX
DR WPI; 2001-639362/73.
DR P-PSDB; ABG10412.
XX
PT New isolated polynucleotide and encoded polypeptides, useful in
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PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity
PS Claim 1; SEQ ID No 10403; 103bp; English.
XX
CC The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AAS64197-AAS94564 represent novel human
CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pcl_sequences.
XX
SQ Sequence 2277 BP; 878 A; 483 C; 473 G; 443 T; 0 other;
XX
Alignment Scores:
Pred. No.: 0.609 Length: 2277
Score: 109.50 Matches: 73
Percent Similarity: 37.21% Conservative: 39
Best local Similarity: 24.25% Mismatches: 107
Query Match: 4.64% Indels: 82
DB: Gaps: 15
US-09-807-459-2 (1-458) x AAS74599 (1-2277)
QY 183 SerPhePheAsnLysLeuSerPheThrGlyLeuPheGlyTyrPglYIleLysArgAla 202
DB 868 ACTTCTCTCCAGAAATTTGGAAAAAACTTAAAGTTCAATATGAAACCAAAAAAGAGCC 927
QY 203 LeuLysGluIleIleArgSerAsnLeu--ProLeuAspIleGlyThrGluHisSerValS 222
DB 928 CTC-----ATCGGCAAGTCAATCCTGAGCCAAAGAAACCAAGCAGACATCAGCGTA 981
QY 222 eraTgLeuGlnHisIleThrSerSerTyrLysAspTyrMetAspThrGlnIleProAlaL 242
DB 982 CCTGACTTCGAACATATCTACAGAGCTACAGTAC-----CAAAAGAGCCTGCTACTGC 1035
QY 242 eupProLysPheAlaLysArgPheSerLeuMetValIaGlnArgLeuLeuIaThrValA 262
DB 1036 TACCAGAAA-----CAGAGATATAGACCAATGGAACAGACAGAGCCCTCAGAAATATG 1089
QY 262 laGlyTyr-----Val-AspThrPro----- 268
DB 1090 CCGCATATCTACAACTATCTGATCTTGACAAACCTCAGAAAAACAAAGCAATGGGGAAG 1149
QY 269 -----TTP---TyrLysLysTrpTyrMetLysLysLysAsnPhmet 281
DB 1150 GATTCCCTATTATTAATAATGCTGCGGAAAAAATGCTAACCAATATGTAAGAAACCTGAAA 1209
QY 282 ValAsnArgValPheIleProThrLysLysPhePheAsnLysGluIleArgGlu----- 299
DB 1210 CTGATCCCTCTTCCCTTACCACTTATACAAAATTAATTCAGATGATGATTAAGACTTACAT 1269
QY 300 -----ProSerLysAlaLeuLysGluLysValSerThrAspThrLysAspLeu 315
DB 1270 GTTAGAGCTAAACCACTAAAACCTTAGAAGAAAAGAACTAGCAATACCATTCAGAC--- 1326
QY 316 PheGluAsnLysIleGlyGlnGlyThrValAspPhePheAsnLysGluIleArgAspPro 335
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Db 1327 -----ATAGGATGGGCAAG---GACTTCATGCTCTAA-----CCA 1359
Oy 336 SerLysAlaLeuLysGluValSerAsnAspAlaLysAspLeuPheGluAsnLys--- 354
Db 1360 CCAAAAGCAACGGCAGCAAAAGCCAAATTTGACAAATGGGATCTTAATTAACCTAAAGAAC 1419
Oy 355 -----IlleGlyGlnGlyThrValAspPheIleAsnAsnGluIleArgAspProSerLys 372
Db 1420 TTCTGCACAGCAAAAGAAATACCATCAGATGAC-----AGGCACTTACAAAA 1470
Oy 373 AlaLeuIleArgLysValSerThrGlyAlaGluAspLeuPheGluAsnLysIleGlyGln 392
Db 1471 -----TGGAGAAATATTTCCACACC----- 1491
Oy 393 GlyThrValAspPheIleAsnAsnGluIleArgAspProSerLysAlaLeuIleArgLys 412
Db 1492 -----TACTCATCTGACAAAGCGCTAAATATCCAGA 1521
Oy 413 ValTyrThrGluAlaAspAspLeuPheGluAsnLysIleGlyGlnGlyThrValAspPhe 432
Db 1522 ATCTACAAATGAACTCCAGCAAAATTTCAAGAAAA----- 1557
Oy 433 IleAsnLysGluIleArgAspProSerLysAlaLeuIleArgLysValSerThrGln 451
Db 1558 ACAAAACAAACCCCATCAAAAAGTGCGGAGACATGACACACATCTTCAAAAAGAA 1614

RESULT 39
AAS79126
ID AAS79126 standard; cDNA; 2277 BP.
XX
XX AAS79126;
XX
XX 13-FEB-2002 (first entry)
XX
XX DNA encoding novel human diagnostic protein #14930.
XX
XX Human; Chromosome mapping; gene mapping; gene therapy; forensic;
XX food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
XX Homo sapiens.
XX
XX WO200175067-A2.
XX
XX 11-OCT-2001.
XX
XX 30-MAR-2001; 2001WO-US08631.
XX
XX 31-MAR-2000; 2000US-0540217.
XX 23-AUG-2000; 2000US-0649167.
XX
XX (HYSE-) HYSEQ INC.
XX
XX Drmanac RT, Liu C, Tang YT;
XX
XX WPI: 2001-639362/73.
XX P-PSDB: AB614939.
XX
XX New isolated polynucleotide and encoded polypeptides, useful in
XX diagnostics, forensics, gene mapping, identification of mutations
XX responsible for genetic disorders or other traits and to assess
XX biodiversity.
XX
XX Claim 1; SEQ ID No 14930; 103pp; English.
XX
XX The invention relates to isolated polynucleotide (I) and
XX polypeptide (II) sequences. (I) is useful as hybridisation probes,
XX polymerase chain reaction (PCR) primers, oligomers, and for chromosome
XX and gene mapping, and in recombinant production of (II). The
XX polynucleotides are also used in diagnostics as expressed sequence tags
XX for identifying expressed genes. (I) is useful in gene therapy techniques
XX to restore normal activity of (II) or to treat disease states involving
XX (II). (II) is useful for generating antibodies against it, detecting or

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CC The polypeptide and polynucleotide sequences have applications in
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CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AS64197-AS94554 represent novel human
CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX
XX Sequence 2277 BP; 878 A; 483 C; 473 G; 443 T; 0 other;

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US-09-807-459-2 (1-458) x AAS79126 (1-2277)
Oy 183 SerPheAsnLysLeuSerPheThrGlyLeuPheGlyTyrGlyIleLysArgAla 202
Db 868 ACTTCTCTCCAGAAATTTGAAAAAACTACTTAAAGTCTATATGAAACCAAAAAGAGCC 927
Oy 203 LeuLysGlnIleIleArgSerAsnLeu--ProLeuAspIleGlyThrGluHisSerVal 222
Db 928 CTC-----ATGCCCAAGCTCAATCTGAGCCAAAGAACCAACCAAGACATCAGCGTA 981
Oy 222 eraArgLeuGlnHisIleThrSerSerTyrLysAspTyrIleTasPthrGlnIleProAla 242
Db 982 CCGACTTCGAACTATACACAAAGGCTACAGTAAC-----CAAAACAGCGTGTACTCG 1035
Oy 242 euProLysPheAlaLysArgPheSerLeuMetValValGlnArgLeuLeuAlaThrVala 262
Db 1036 TACCAAAA-----CAGAGTATAGACCAATGGAACAGAACAGACCCCTCAGAAATATG 1089
Oy 262 IAGlyTyr-----Val-AspThrPro----- 268
Db 1090 CCGCATATCTACAACTATCTGATCTTGCAAAACCTCACAAAACCAACATGGGGAAG 1149
Oy 269 -----Tyr--TyrLysLysTyrTyrIleLysLeuLysAsnPheMet 281
Db 1150 GATTCCTATTATTAATAATGTCTGCGAATAATGCTTACCATATGTAGAAAGCTGA 1209
Oy 282 ValAsnArgValPheIleProThrLysLysPheAsnLysGluIleArgGlu----- 299
Db 1210 CTGGATCCCTCTTACACCTTATACAAAATTAATTAAGATGAAGACTTACAT 1269
Oy 300 -----ProSerLysAlaLeuLysGluLysValSerThrAspThrLysAspLeu 315
Db 1270 GTTAGAGCTAAACCATTAACCCCTAGAGAAAACCTAGACATACCATTCAGGAC--- 1326
Oy 316 PheGluAsnLysIleGlyGlnGlyThrValAspPheAsnLysGluIleArgAspPro 335
Db 1327 -----ATAGGATGGGCAAG---GACTTCATGCTCTAA-----CCA 1359
Oy 336 SerLysAlaLeuLysGluValSerAsnAspAlaLysAspLeuPheGluAsnLys--- 354
Db 1360 CCAAAAGCAACGGCAGCAAAAGCCAAATTTGACAAATGGGATCTTAATTAACCTAAAGAAC 1419
Oy 355 -----IlleGlyGlnGlyThrValAspPheIleAsnAsnGluIleArgAspProSerLys 372
Db 1420 TTCTGCACAGCAAAAGAAATACCATCAGATGAC-----AGGCACTTACAAAA 1470
Oy 373 AlaLeuIleArgLysValSerThrGlyAlaGluAspLeuPheGluAsnLysIleGlyGln 392
Db 1471 -----TGGAGAAATATTTCCACACC----- 1491

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Job time : 1278 secs

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